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(84) Oligosaccharides as a hair growth promotor.

(87) Novel esterified oligosaccharides including at least one esterified disaccharide unit consisting of a uronic acid residue and a hexosamine residue can be topically applied to mammalian skin to improve the appearance of hair and other benefits. These oligosaccharides can be synthesised by chemical cleavage of an intact glycosaminoglycan chain or by condensation of hexosamine and uronic acid residues.

**EP 0 211 610 A2**

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## TITLE MODIFIED

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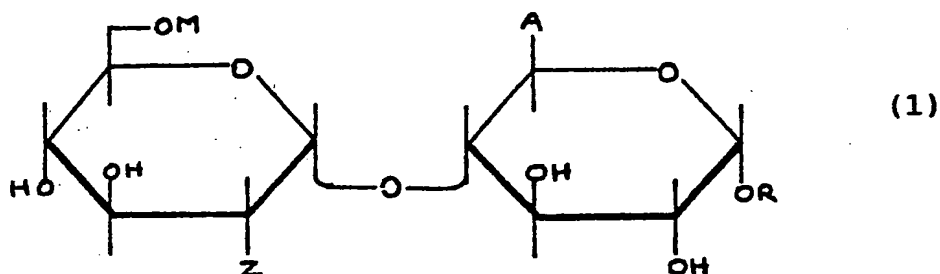
J.3023/1

OLIGOSACCHARIDESField of invention

- 5       The invention relates to novel oligosaccharides, particularly to esterified disaccharides containing a uronic acid residue and a hexosamine residue joined through a glycosidic linkage. The invention also relates to the synthesis of these esterified disaccharides and to their use, particularly in cosmetic and pharmaceutical compositions.
- 10

Prior Art

- 15       Certain  $\alpha$ -1,4 esterified disaccharides, together with their synthesis and use in controlling the coagulation of blood, are described by Choay S.A. in EP-A-0 064 012. These  $\alpha$ -1,4 disaccharides are said to have the structure:



10 where Z represents a functional nitrogen group, such as an azide or a group having the structure -NHB, in which B represents -H or a functional group such as acetyl or sulphate as a salt with an organic or mineral cation;

15 M represents -H or  $-\text{SO}_3\text{M}_1$ , where  $\text{M}_1$  is an organic or metallic cation, particularly an alkali metal; or an acetyl group;

20 R represents a  $\text{C}_1$  to  $\text{C}_4$  alkyl radical, especially methyl; or an aryl radical;

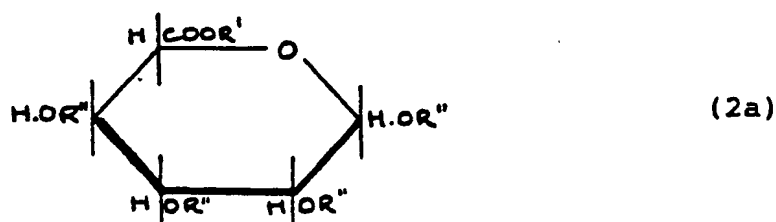
25 A represents a functional group such as an acid or  $-\text{COOR}_1$ , where  $\text{R}_1$  represents -H or a  $\text{C}_1$  to  $\text{C}_4$  alkyl radical, especially methyl; or a metal, especially an alkali metal.

#### COMPOUNDS PER SE

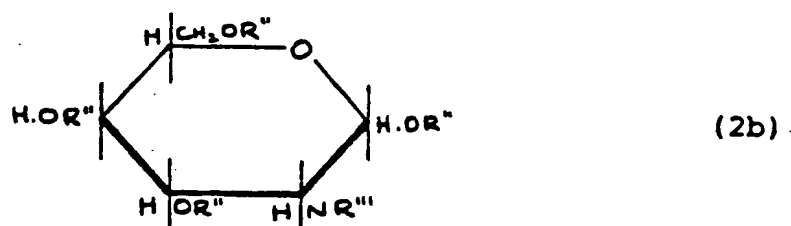
30 The novel esterified oligosaccharides of the invention differ from the disaccharides described by Choay S.A.

Definition of the invention

Accordingly, the invention provides esterified oligosaccharides including at least one esterified disaccharide unit consisting of a uronic acid residue having the structure:



15 and a hexosamine residue having the structure:



25 where R' is C<sub>3</sub> to C<sub>10</sub> alkyl or  $\text{COOR''}$   
 R'' is -H, C<sub>1</sub> to C<sub>4</sub> alkyl, -CO(CH<sub>2</sub>)<sub>m</sub>CH<sub>3</sub>, -SO<sub>3</sub>M,  
 R''' is -H, -CO(CH<sub>2</sub>)<sub>m</sub>CH<sub>3</sub>, or -SO<sub>3</sub>M,  
 M is -H, or a metallic or organic cation  
 n is 0 or an integer of from 1 to 7, and  
 m is 0 or the integer 1 or 2;

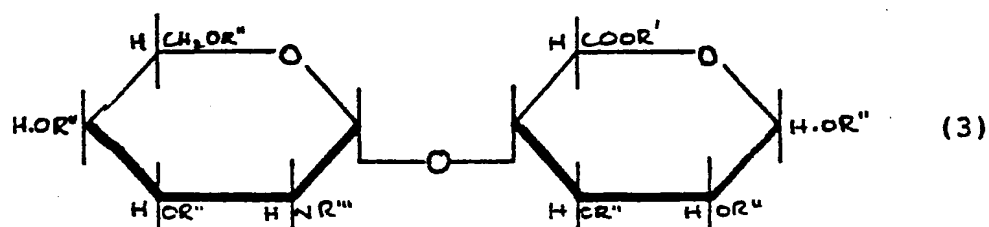
30 the groups designated R'' being the same or different, one R'' group from each pyranose ring structure being linked by a glycosidic linkage having the configuration α-1,3; α-1,4; β-1,3 or β-1,4; and the -COOR', -CH<sub>2</sub>OR'' and -OR'' groups being of either configuration with respect to the pyranose rings;

35

provided that when the esterified oligosaccharide is an esterified disaccharide unit having the structure:

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10



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then R' is C<sub>5</sub> to C<sub>10</sub> alkyl or  $\begin{array}{c} \text{COOR''} \\ | \\ -\text{CH}(\text{CH}_2)_n\text{CH}_3 \end{array}$

20

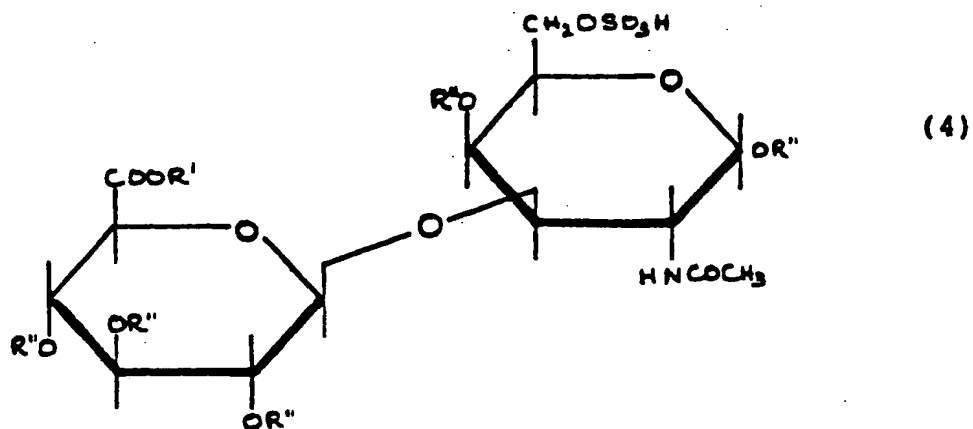
### The esterified oligosaccharide

The esterified oligosaccharide according to the invention can comprise from 2 to 8, preferably from 2 to 4 saccharide residues, of which at least two saccharide residues form the esterified disaccharide unit as herein defined.

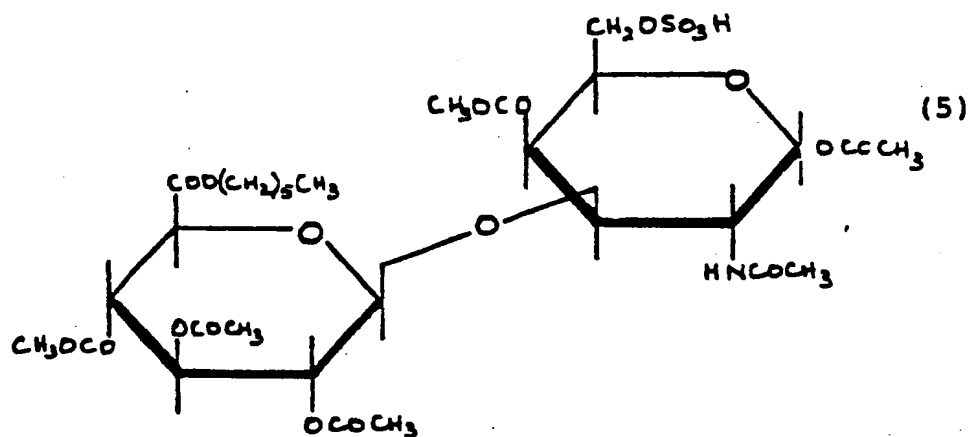
Preferably, the esterified oligosaccharide is an esterified disaccharide consisting of a uronic acid residue having the structure (2a) linked via a glycosidic linkage to a hexosamine residue having the structure (2b).

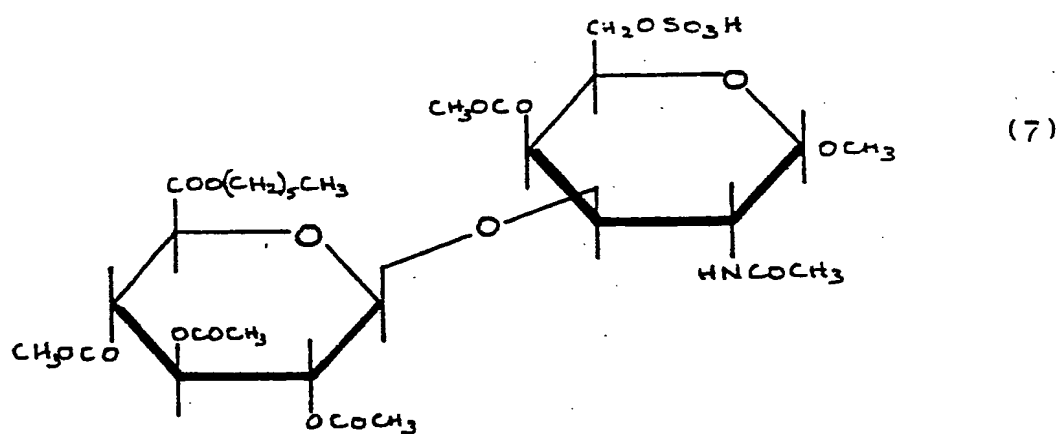
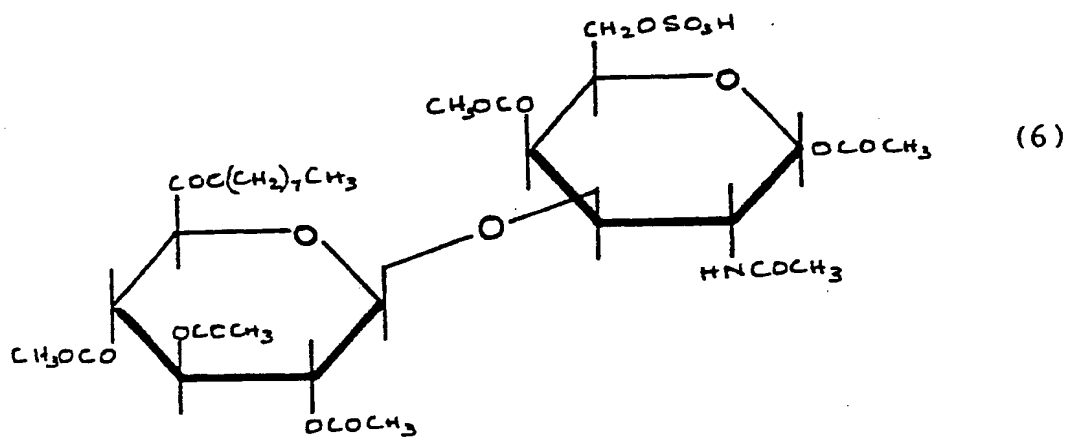
Particularly preferred examples of the esterified disaccharides of the invention have the following structures, where the designation of R', R'', R''' and are as hereinbefore defined.

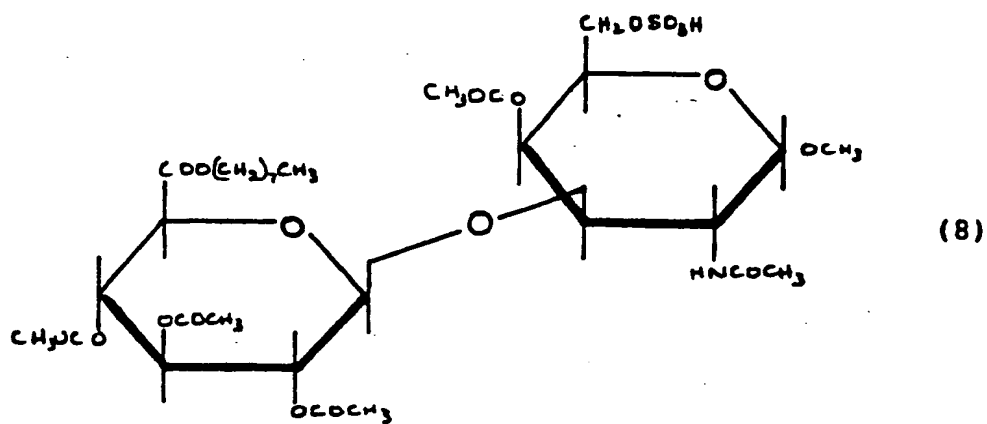
A first generic structure is:



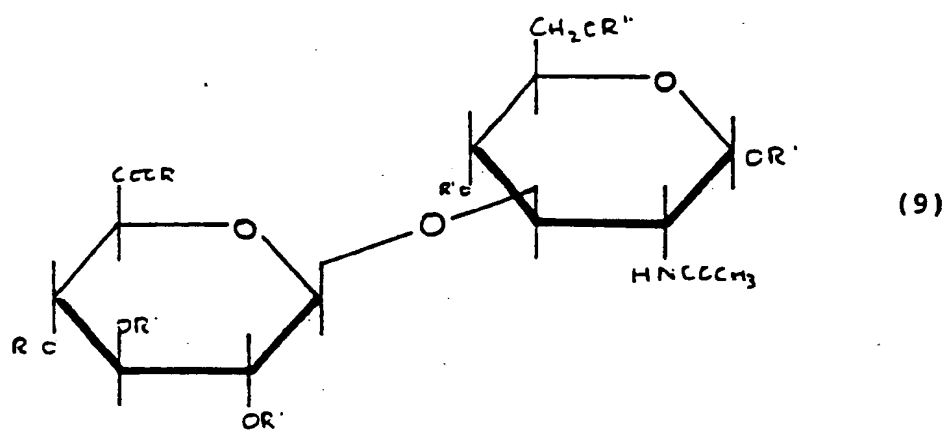
Specific examples of the esterified disaccharide (4) have the structures shown as (5), (6), (7) and (8).





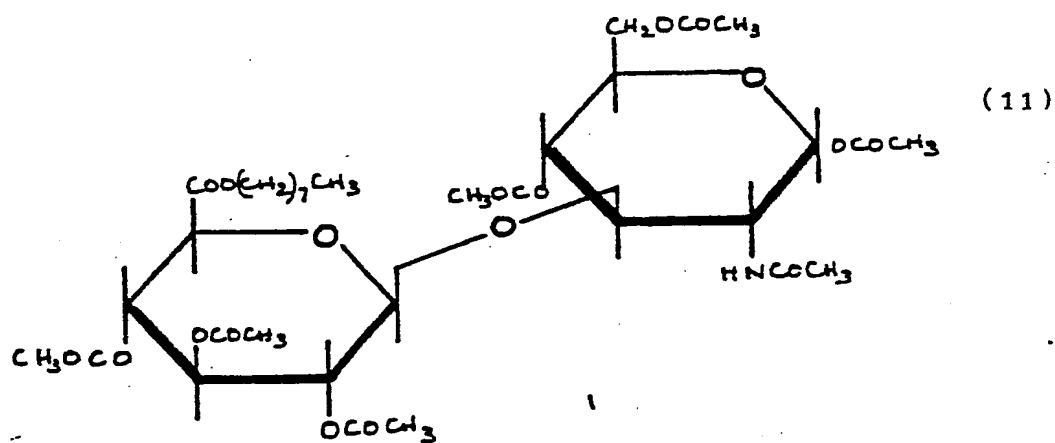
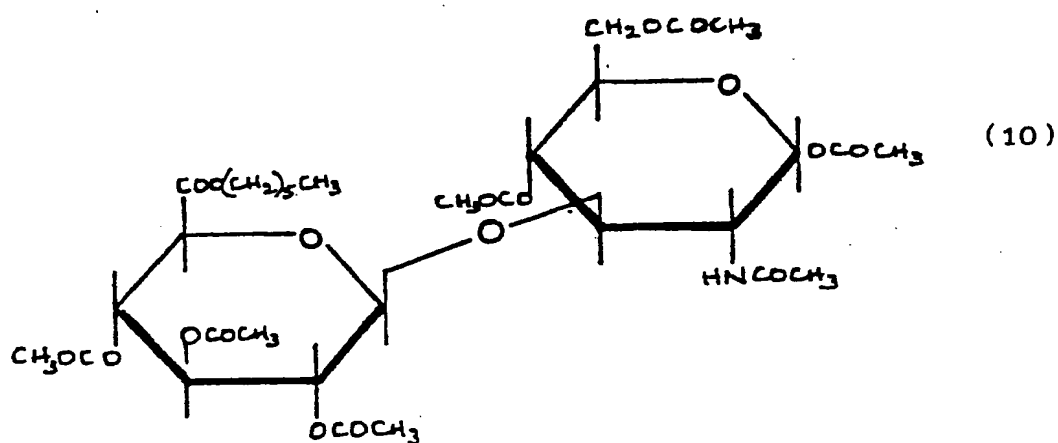


A further generic structure is:

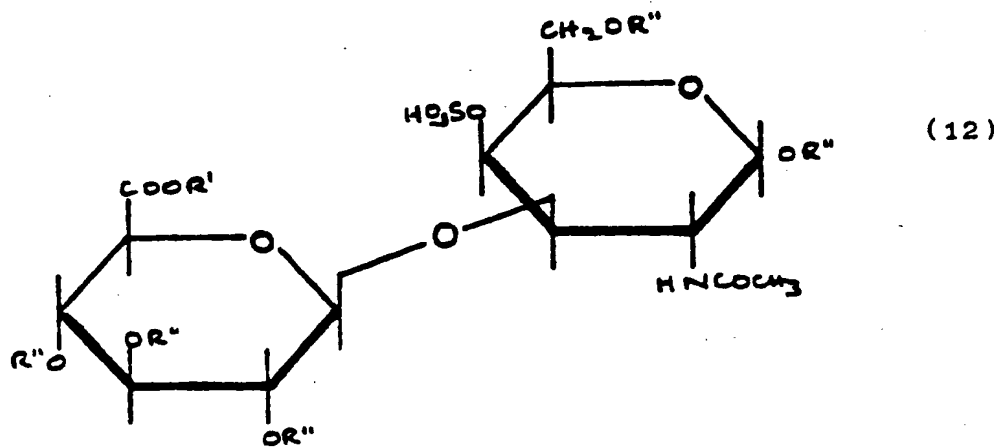




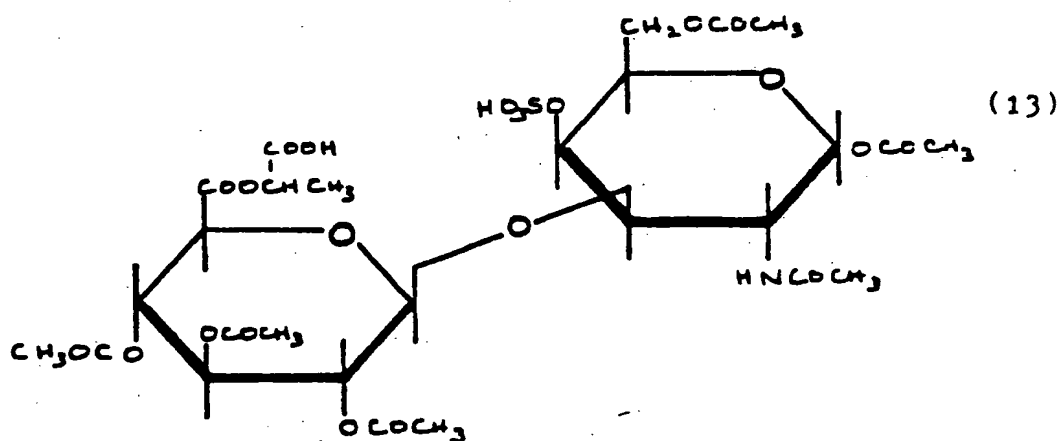
Specific examples of the esterified disaccharide (9) have the structures shown in (10) and (11).



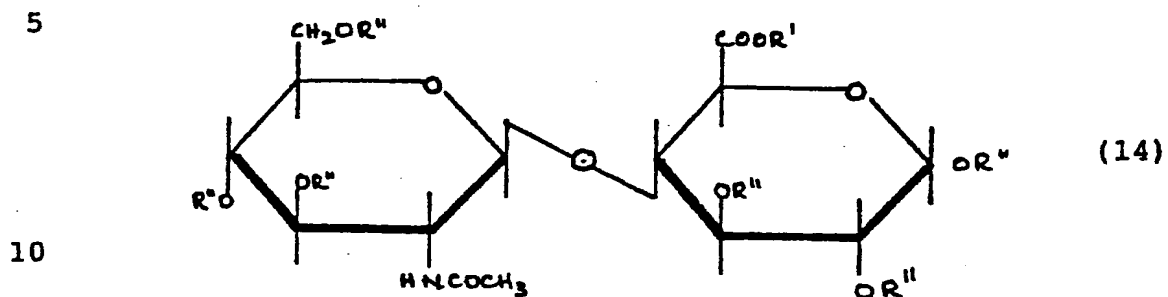
A further generic structure is:



A specific example of the esterified disaccharide (12) has the structure shown in (13).



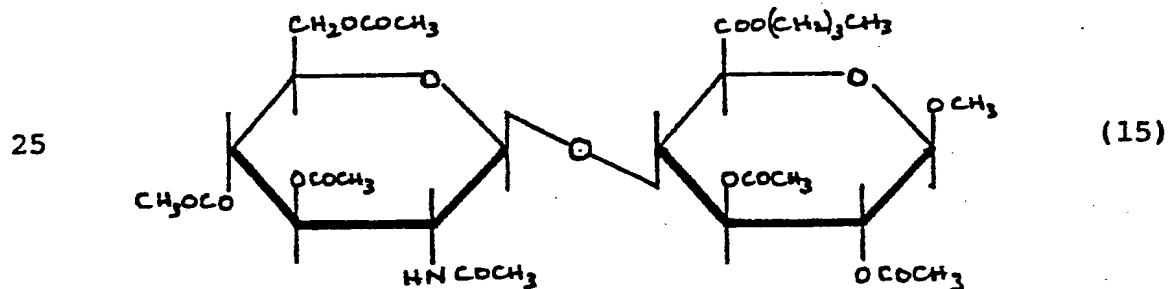
A further generic structure is:



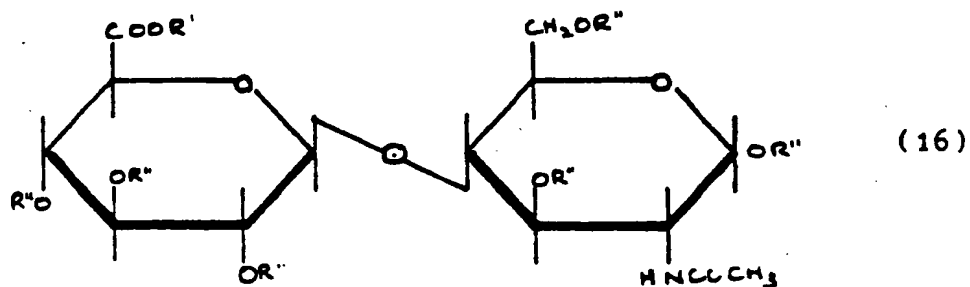
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A specific example of the esterified disaccharides (14) has the structure shown in (15)

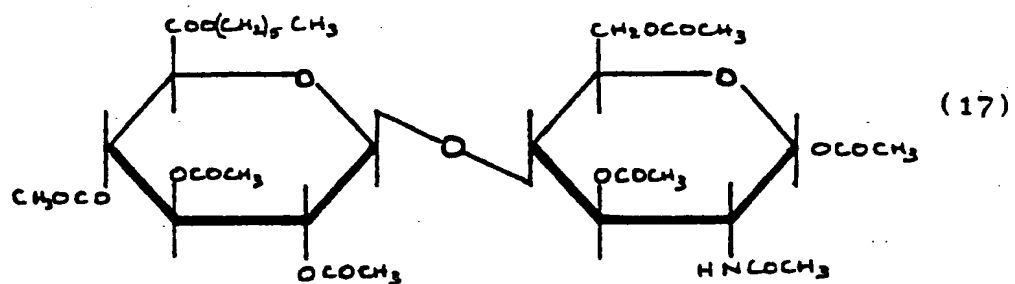
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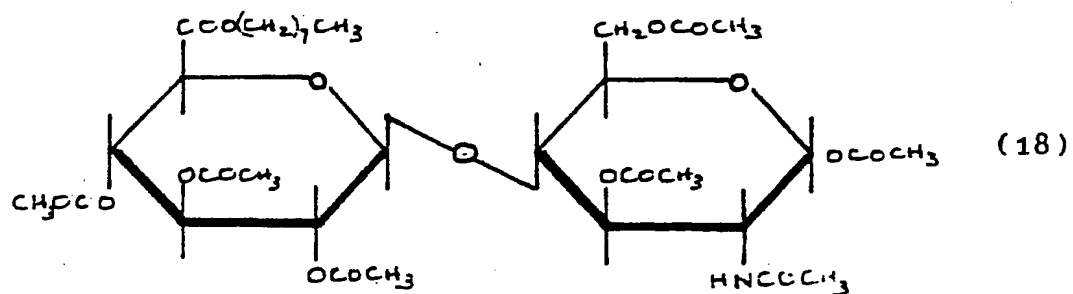


A further generic structure is:

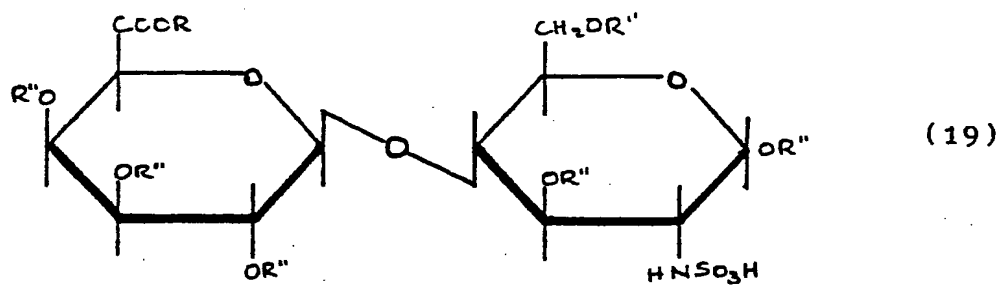


Specific examples of the esterified disaccharide (16) have the structures shown in (17) and (18).

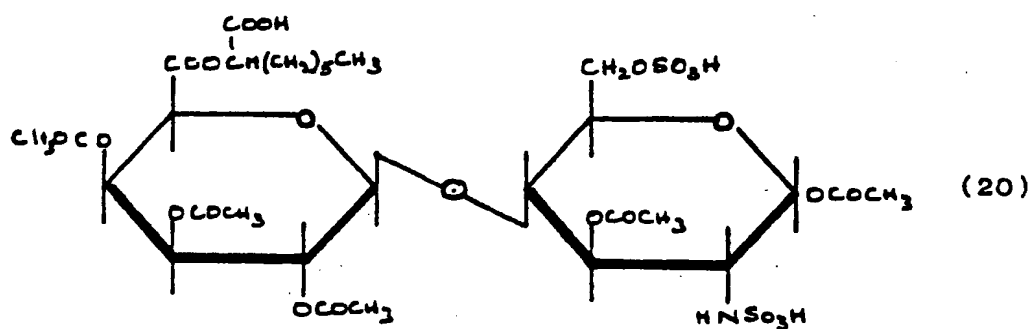




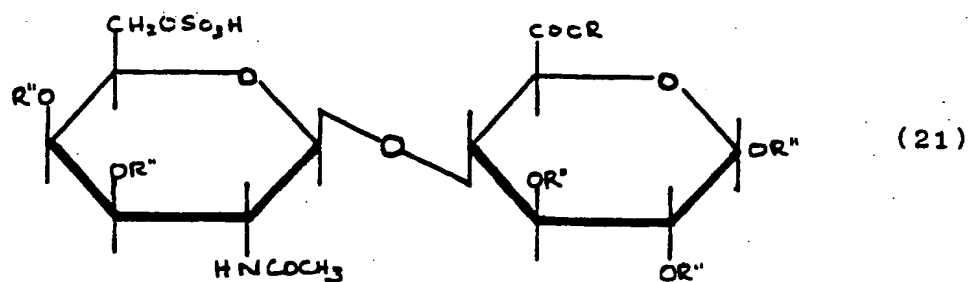
A further generic structure is:



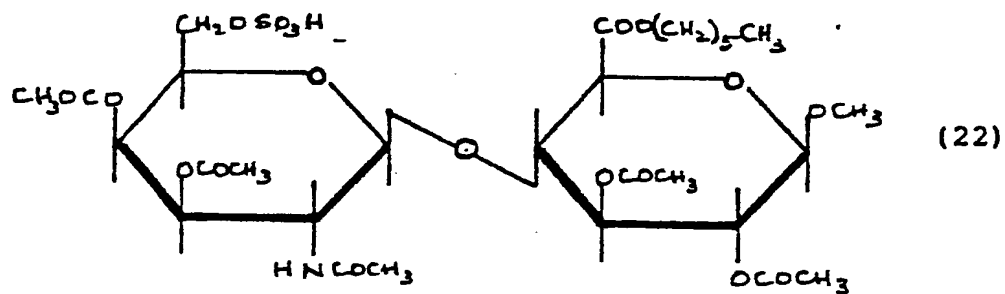
A specific example of the esterified disaccharide (19) has the structure shown in (20).



A further generic structure is:

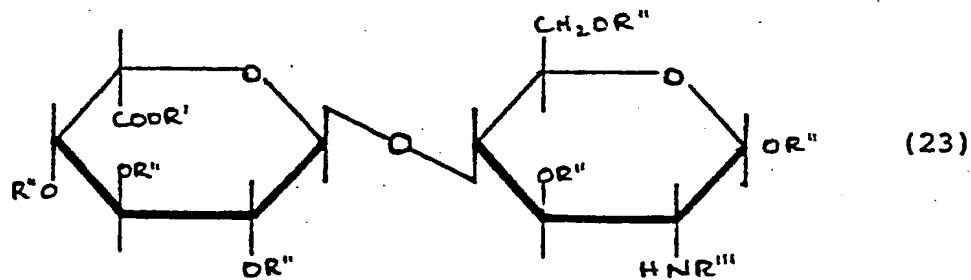


A specific example of the esterified disaccharide (21) has the structure shown in (22):

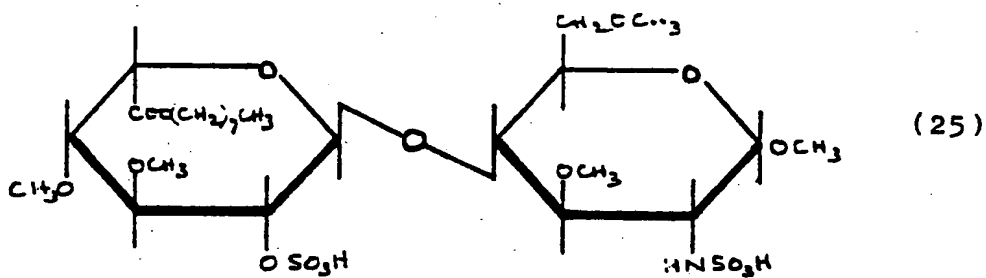
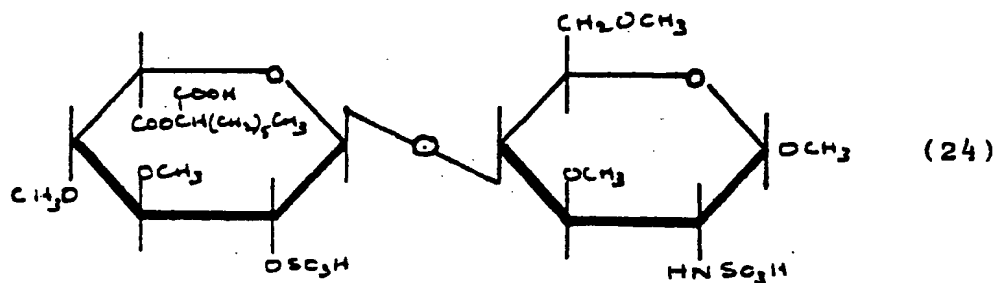


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A further generic structure is:

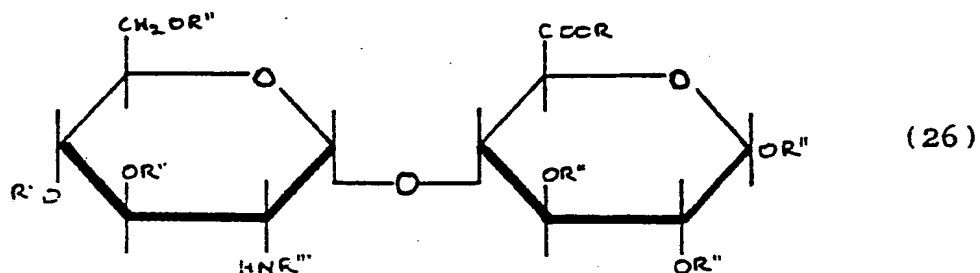


Specific examples of the esterified disaccharide (23) has the structures shown in (24) and (25).

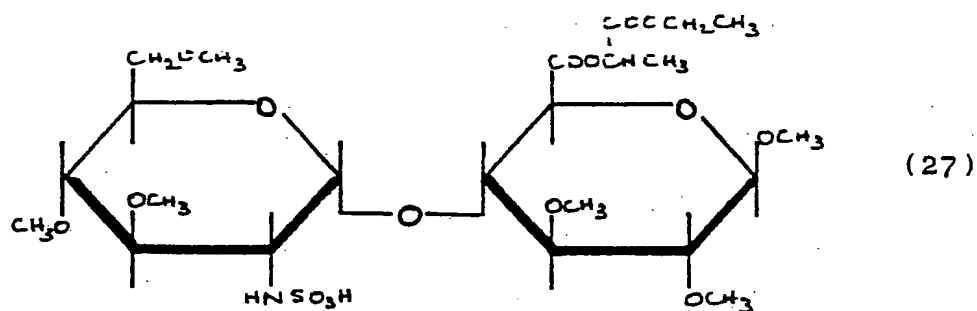




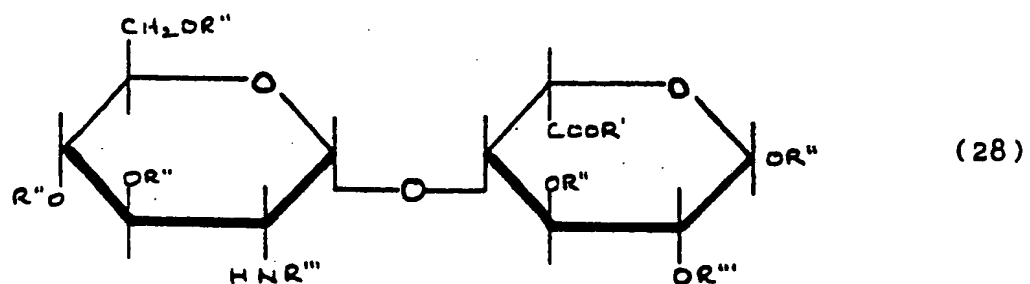
A further generic structure is:



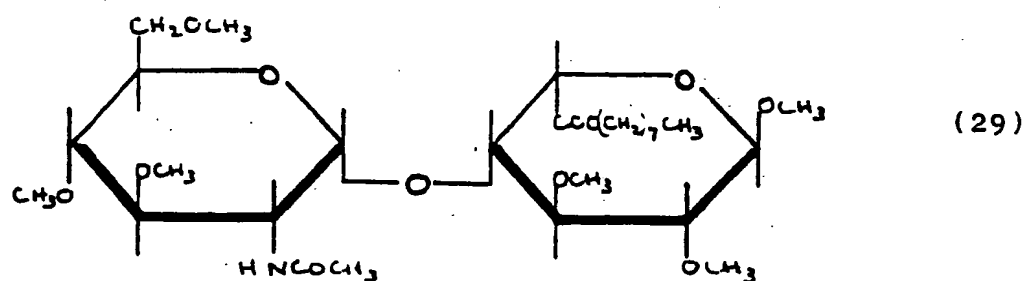
A specific example of the esterified disaccharide (26) has the structure shown in (27).



A further generic structure is:



A specific example of the esterified disaccharide (28) has the structure shown in (29).



SYNTHESIS OF THE NOVEL ESTERIFIED OLIGOSACCHARIDES

Two possible routes for synthesising the novel esterified oligosaccharides according to the invention are  
5 envisaged.

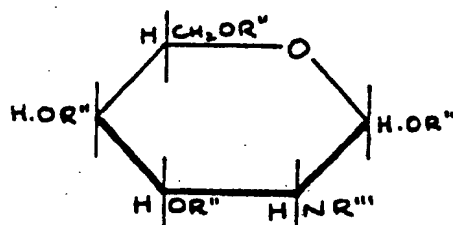
1. Hydrolysis of polysaccharides

Intact glycosaminoglycan chains can be subjected to  
10 partial hydrolysis to yield oligosaccharides comprising at least one disaccharide having a uronic acid residue linked to a hexosamine residue. These oligosaccharides can then be modified to provide the corresponding esterified oligosaccharides of the invention. This synthesis is  
15 particularly suited to the production of esterified oligosaccharides having the structures of the generic types (4), (9) and (12).

Definition of the invention

Accordingly, the invention provides a process for the synthesis of an esterified oligosaccharide having a generic structure chosen from structures (4), (9) and (12), which comprises the steps of:

( i) Subjecting a glycosaminoglycan chain to chemical cleavage to yield oligosaccharide fragments containing at least one disaccharide unit comprising a hexosamine residue having the structure:

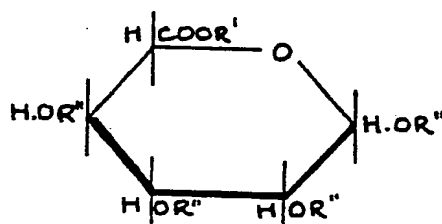


(2b)

which is glycosidically linked to the C-1 position of a uronic acid residue having the structure:

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(2a);  
and

15

( ii) modifying the oligosaccharide by at least one process step chose from:

acylation of free hydroxyl groups,  
acylation of free amino groups,  
sulphation of free hydroxyl groups,  
sulphation of free amino groups,  
esterification of free hydroxyl groups, and  
etherification of free hydroxyl groups,

25

to provide a structure of the generic type chosen from structures (4), (9) and (12).

Preferred glycosaminoglycan chains which form the starting materials for this synthesis are chondroitin sulphate and hyaluronic acid.

Of the modifying steps which can be applied to the oligosaccharide in order to obtain the esterified oligosaccharide of the invention, it is preferred that these are carried out in a stepwise fashion to acylate the

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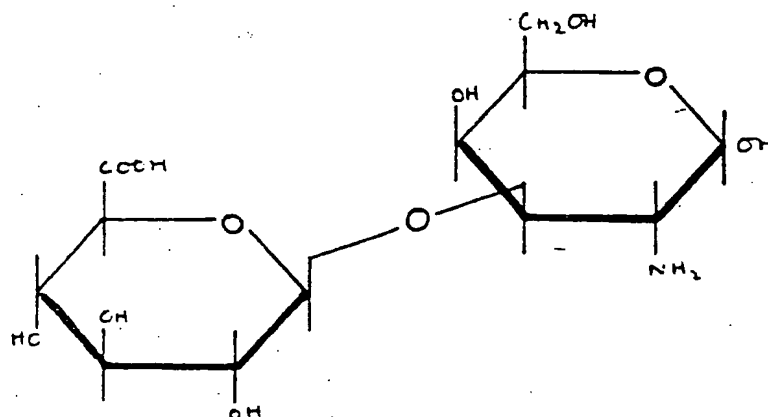
2-amino group of the hexosamine moiety to esterify or acylate the remaining free hydroxyl groups and to esterify the carboxyl group on the uronic acid moiety.

- 5           Chemical cleavage of the intact glycosaminoglycan chains is preferably achieved by acid hydrolysis or enzymic digestion.

10           In order to illustrate Method 1, the synthesis of the esterified disaccharide having the structure (5) starting from chondroitin sulphate is now described.

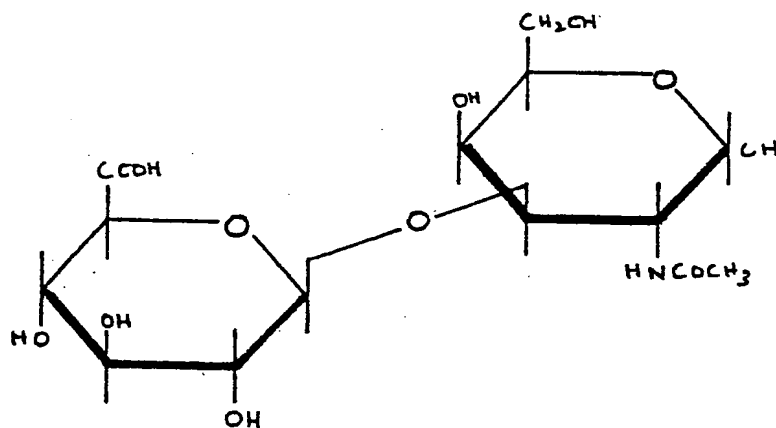
15           The disaccharide chondrosine (30) was obtained from chondroitin sulphate by acid hydrolysis. 10g of chondroitin sulphate was hydrolysed with 200ml of 2N sulphuric acid by refluxing at 100°C for 4 hours. The resulting hydrolysate was chromatographed on Dowex 50W-X8 (200-400 mesh) resin in the hydrogen form. The column was eluted with 0.2N sulphuric acid and all fractions were tested with ninhydrin/ferric chloride reagent. The first large, broad peak gave a yield of chondrosine after neutralisation with saturated barium hydroxide and lyophilisation to a dry powder.

25           The method employed was that described by Olavesen & Davidson, J.Biol.Chem. 240(3), 992-996 (1965).



(30)

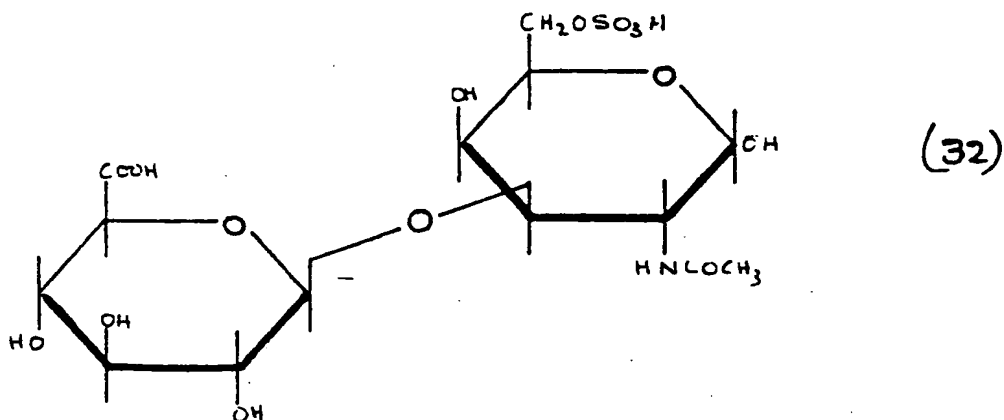
- N-acetylchondrosine (31) was prepared by selective N-acetylation of chondrosine using acetic anhydride. 500mg chondrosine was dissolved in 500 ml of bicarbonate buffer and cooled to 0-5°C. (Bicarbonate buffer was prepared by mixing 10ml 0.1M  $\text{Na}_2\text{CO}_3$  and 90ml 0.1M  $\text{NaHCO}_3$  having a pH value of 9. To this was added 0.22ml of acetic anhydride with stirring. The mixture was then left standing at 0°C overnight. It was then added to a Dowex 50W-X8 (200-400) column in the acid form. Elution with three bed volumes of distilled water and lyophilisation of the aqueous eluate produced a white fluffy solid which was characterised as N-acetylchondrosine by I.R. and N.M.R. spectroscopy.
- The method employed was that described by Olavesen & Davidson, Ibid.



(31)

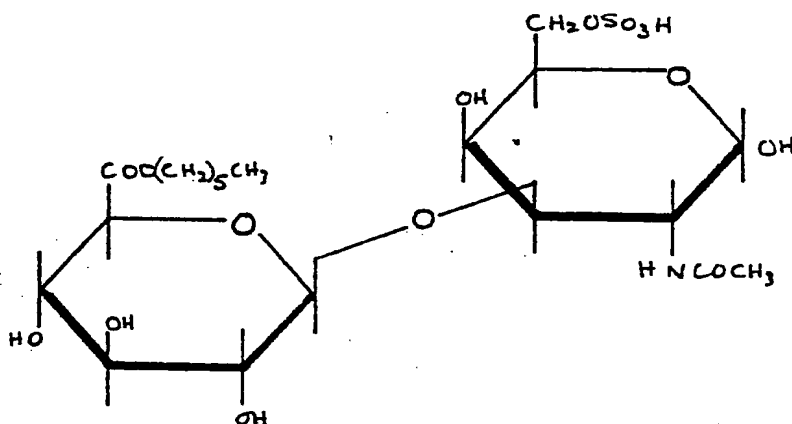
Sulphation of the product in the C-6 position of the hexosamine moiety was achieved by using triethylamine-N-sulphonic acid. To a mixture of 5ml dimethylformamide, 7.5ml benzene and 0.5ml pyridine was added 500mg of N-acetylchondrosine after it had been evaporated several times from dry benzene. 9.12mg of triethylamine-N-sulphonic acid was added and the reaction vessel shaken for 26 hours at room temperature. The resulting solution was evaporated at reduced pressure to a yellow syrup (bath temp. <50°C) which was extracted with 2 x 5ml ether. The resulting oil was extracted with 10ml chloroform to remove any residual sulphonic acid. Chloroform washings and insoluble solids were washed in situ with 4 x 5ml water. The water fractions were combined and lyophilised to give N-acetylchondrosine-6-O-sulphate (32), as a white fluffy solid which was characterised by I.R. and N.M.R. and by T.L.C.

The method employed was that described by Olavesen & Davidson, Biochim. Biophys. Acta., 101, 245-251 (1965).





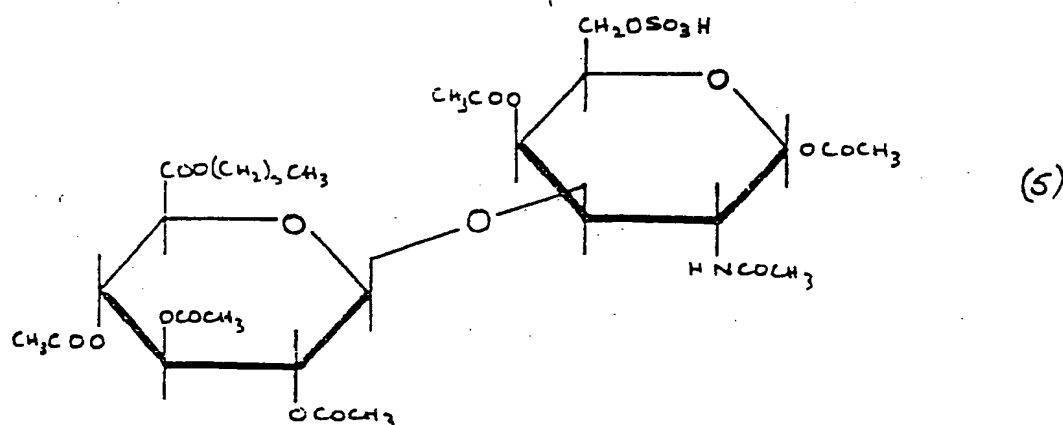
The N-acetyl chondrosine-6-O-sulphate (32) was esterified using a modified Pummerer reaction. 500mg N-acetylchondrosine 6-O-sulphate was added to a large round bottomed flask containing 5ml of hexanol and 3.4g of  
5 Amberlite IR-120 resin in the hydrogen form. The mixture was refluxed at 100°C for 4 hours and then the reaction was forced to completion by distillation of the hexanol and formed water. The residual resin was removed by  
10 filtration and washing with water. The product having the structure (33) was obtained by lyophilisation and was characterised by I.R., N.M.R. and T.L.C.



(33)

The fully acetylated disaccharide (5) was prepared using the acetic anhydride/pyridine reagent. 500mg N-acetylchondrosine-6-O-sulphate-C<sub>6</sub> ester was dissolved in 30ml dry pyridine and to this was added 20ml acetic anhydride. The mixture was then cooled to 0°C and stirred until all the disaccharide had dissolved. It was then stirred at room temperature for 48 hours. The resulting brown solution was poured into 300ml of iced water which was extracted with 3 x 50ml chloroform. Chloroform solution was evaporated to dryness under reduced pressure at 45°C to give a yellow syrup. This was treated with ethanol to give an amorphous solid which was subsequently eluted with ethyl acetate from a silicic acid column and lyophilised to produce an amorphous white powder.

The method employed was that employed by Olavesen & Davidson, J.Biol.Chem. 240(3), 992-996 (1965).



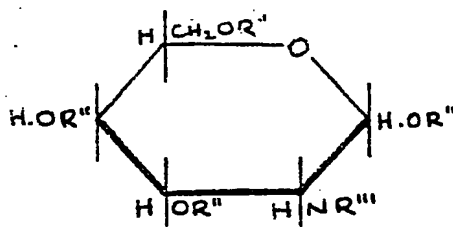
2. Condensation of uronic acid and hexosamine residues

Esterified oligosaccharides according to the invention having at least one uronic acid residue linked in the  $\beta$  configuration through the C-1 position to a hexosamine residue, or certain esterified oligosaccharides according to the invention having at least one hexosamine residue linked in the  $\alpha$  or  $\beta$  configuration through the C-1 position to a uronic acid residue, can be prepared by condensation of relevant uronic acid and hexosamine residues. This synthesis is particularly suited to the production of esterified oligosaccharides having the structures of the generic types (14), (16), (19), (21), (23), (26) and (28).

Definition of the invention

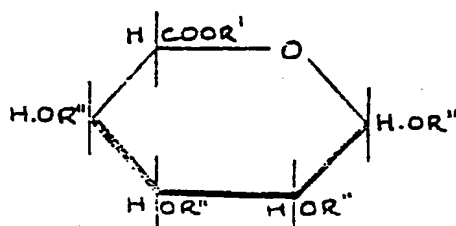
Accordingly, the invention also provides a process for the synthesis of an esterified oligosaccharide having at least one uronic acid residue linked in the  $\beta$  configuration through the C-1 position to a hexosamine residue, or having at least one hexosamine residue linked in the  $\alpha$  or  $\beta$  configuration through the C-1 position to a uronic acid residue, which process comprises the steps of:

( i) condensing a hexosamine residue having the structure:



(2b)

with a uronic acid residue having the structure:



(2a)

to form an oligosaccharide intermediate, the functional groups of which residues have been modified or protected in such a way that glycosidic bond formation can only occur between the C-1 position of one residue and a free hydroxyl group in the C-3 or C-4 position of the other residue, the  $\alpha$  or  $\beta$  configuration of the glycosidic linkage so formed being dictated by the nature of the leaving group on the C-1 position; and

( ii) removal of at least one protecting group from the oligosaccharide intermediate by one or more process steps in any sequence chosen from:

reduction,  
acid catalysis, and  
base catalysis;

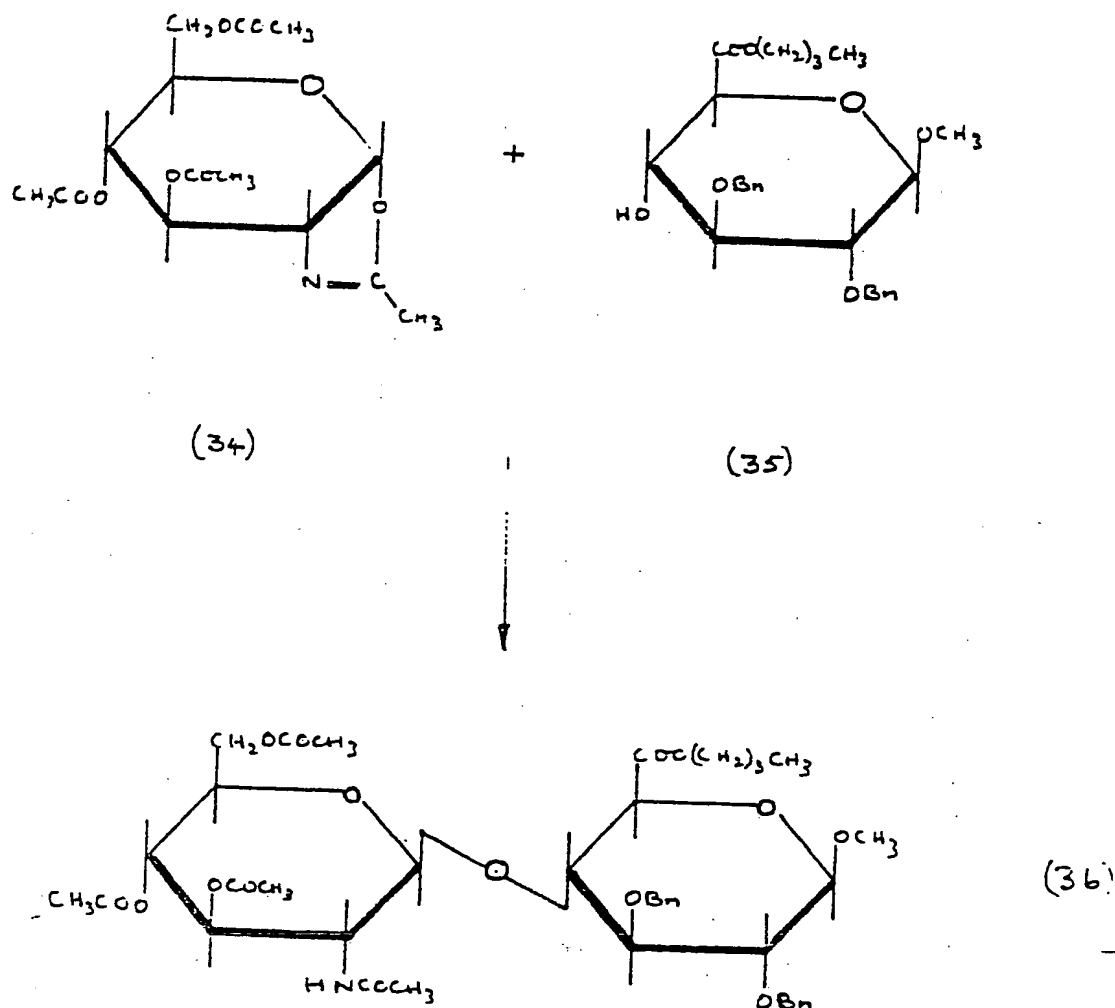
to provide the esterified oligosaccharide.

It is preferred that this process further comprises modifying the oligosaccharide intermediate by one or more process steps in any sequence, chosen from:

acylation of free hydroxyl groups  
acylation of free amino groups  
sulphation of free amino groups  
esterification of free hydroxyl groups, and  
5 etherification of free hydroxyl groups.

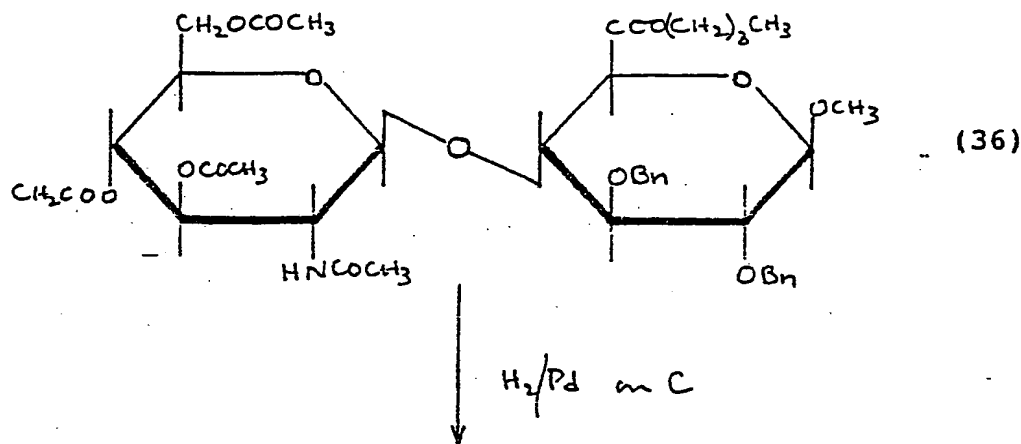
In order to illustrate Method 2, the synthesis of the esterified disaccharide having the structure (15) by condensation of monosaccharides is now described.

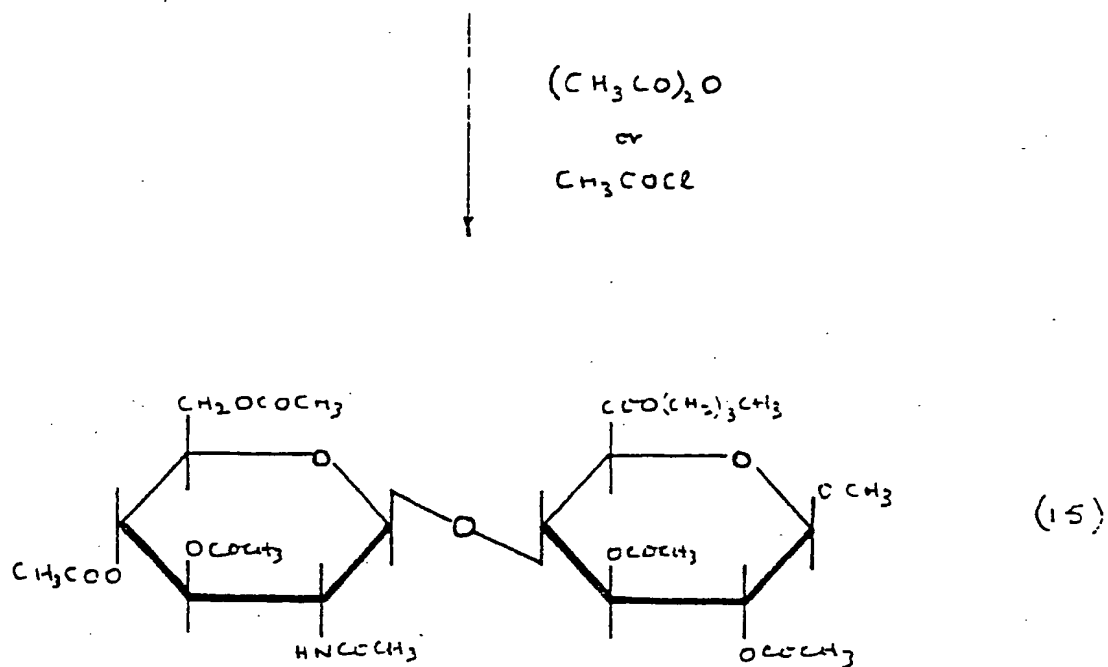
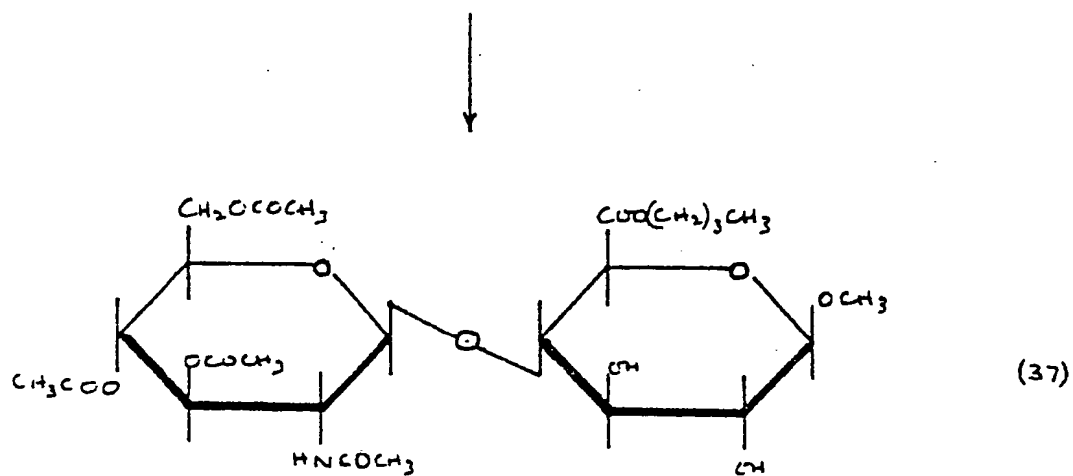
- 5 The esterified disaccharide having the structure (15) was prepared by condensation of monosaccharides (34) and (35), as shown below, to give the ( $\beta$ -1,4) disaccharide (36):



The preparation of (34), the oxazoline derivative of N-acetylglucosamine, was by reaction of N-acetylglucosamine with acetyl chloride by the method described by Horton & Wolfrom in J.Org.Chem. 27, 1794 (1962), to insert an  $\alpha$ -Cl at the C-1 position, followed by oxazoline formation in the presence of tetraethylammonium chloride by the method described by Lemieux & Driguez J.Am.Chem.Soc. 97, 4063 (1975). The oxazoline was a syrup having an  $[\alpha]_D$  of  $+10^\circ$ . Compound (35) was synthesised by esterification, with n-butanol in the presence of sulphuric acid, of the sodium salt of the corresponding uronic acid. The latter was prepared as described in EP-A-O 084 999 and EP-A-O 064 012.

Condensation of (34) and (35) was carried out in the presence of nitromethane and p-toluenesulphonic acid as described by David & Veyrieres in Carbohydrate Research 40, 23 (1975). This paper also provided the method for the reduction of (36) by hydrogenation using palladium on carbon as catalyst, to give compound (37):







The final step, yielding disaccharide (15) was acetylation of the remaining two ring hydroxyl groups on structure (37) by conventional methods using either acetic anhydride or acetylchloride.

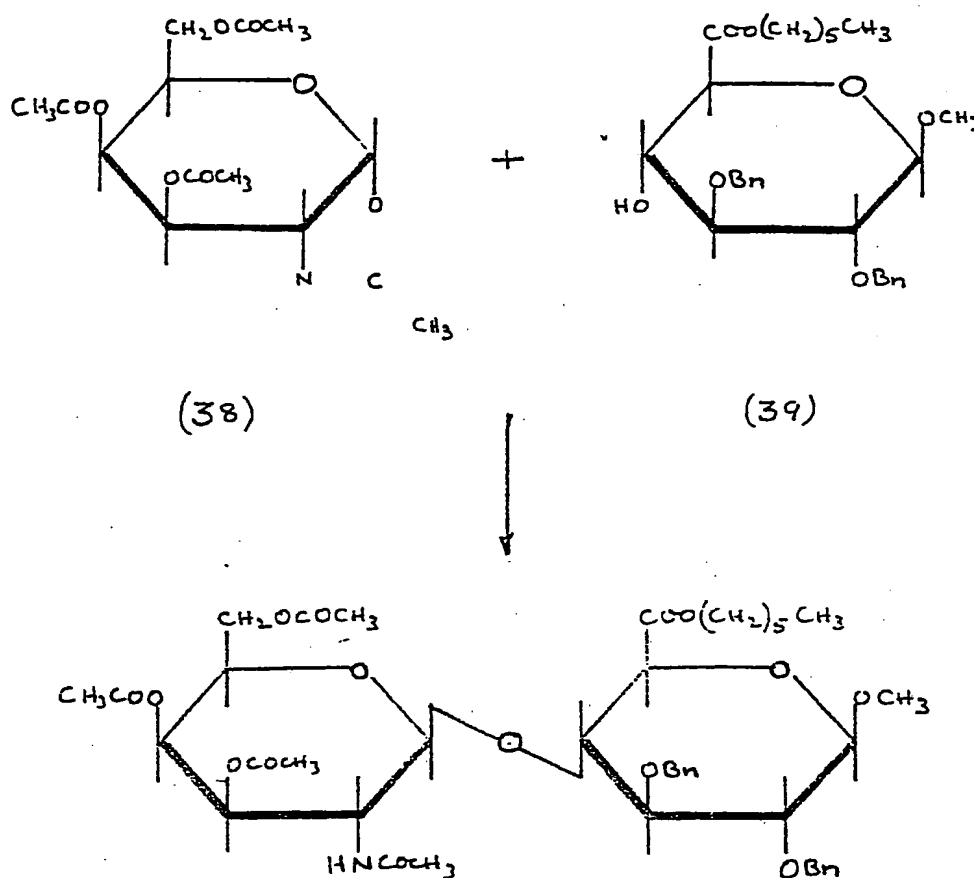
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Method 2 can also be illustrated with reference to the synthesis of the esterified disaccharide (22) by condensation of monosaccharides as follows.

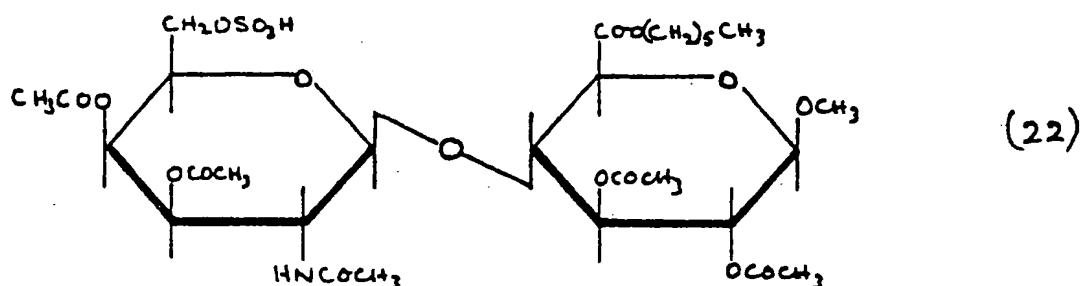
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A similar sequence of reactions as described for disaccharides (15) was followed to prepare disaccharide (22) except that in this case the oxazoline derivative of N-acetyl galactosamine (38) was prepared and condensed with the hexyl ester (39) shown below:

15



Disaccharide (40) was deacetylated with sodium methoxide and then reduced as described in the synthesis of disaccharide (5) above to remove the benzyl protecting groups. The resulting disaccharide was sulphated in the C-6 position by the method described above for disaccharide(5). Finally, remaining free hydroxyl groups were acetylated to yield disaccharide (22):



COSMETIC AND PHARMACEUTICAL COMPOSITIONS CONTAINING  
ESTERIFIED OLIGOSACCHARIDES

5       The invention also relates to cosmetic and  
pharmaceutical compositions for topical application to  
mammalian skin containing novel esterified  
oligosaccharides, particularly esterified disaccharides,  
as hereinbefore described, and other esterified  
disaccharides such as those disclosed by Choay S.A. in  
10   EP-A-0 064 012. These compositions are particularly  
useful in promoting or enhancing the growth of hair, more  
particularly on the human scalp.

The Hair Growth Cycle

15

It should be explained that in most mammals, hair  
does not grow continuously, but undergoes a cycle of  
activity involving alternate periods of growth and rest.  
The hair growth cycle can be divided into three main  
20   stages, namely:

i)    the growth phase known as anagen, during which the  
hair follicle penetrates deep into the dermis with  
the cells of the bulb dividing rapidly and  
25    differentiating to form the hair,

ii)   the transitional stage known as catagen which is  
heralded by the cessation of mitosis, and during  
which the follicle regresses upwards through the  
30    dermis and hair growth ceases,

iii)  the resting stage known as telogen in which the  
regressed follicle contains a small secondary germ  
with an underlying ball of tightly packed dermal  
35    papilla cells.

The initiation of a new anagen phase is revealed by rapid cell proliferation in the germ, expansion of the dermal papilla and elaboration of basement membrane components. The hair cycle is then repeated many times  
5 until, most of the hair follicles spend an increasing proportion of their time in the telogen stage, and the hairs produced become finer, shorter and less visible; this is known as terminal to vellus transformation. In the human male subject this consequence is seen as the onset  
10 of male pattern baldness.

#### Alleged Baldness Cures

Although there have been many claims in the  
15 scientific literature to the promotion or maintenance of hair growth or regrowth, by the topical application of hair tonics and the like, particularly in the human male subject, none has yet been widely accepted by the consumer as being both safe and effective. Perhaps the  
20 only means which has met with partial success for growing hair on the bald or balding human head is by transplantation of hair to the bald areas. This is however an extremely painful operation and is not always successful. Furthermore, it is immediately apparent to  
25 the casual observer that the subject has received a hair transplant, and it may take many months or even years before hair regrowth, following this operation, assumes an appearance which resembles that of the original naturally growing hair.

30

Among the many hair regrowth studies that have been reported in the literature, the work of Meyer et al (1961) in the Proceedings of the Society of Experimental and Biological Medicine, 108, 59-63, is worthy of mention.  
35 Meyer and his co-workers repeatedly injected acid mucopolysaccharides intracutaneously into the skin of

shaved rabbits and reportedly observed initiation of the hair growth cycle with stimulation of hair growth which in some instances appeared to be thicker than usual. They found that heparitin (also known as heparan) sulphate was particularly active, while dermatan sulphate and chondroitin-6-sulphate were also active in this respect, but to a lesser extent.

It has also been reported by Frajdenrajch in EP-A-0 035 919 to include chondroitin sulphate in a hair composition in order to prevent loss and encourage growth of the hair.

Also, Shansho Seijaku in JA-59/186911 describe a shampoo containing a mucopolysaccharide such as chondroitin sulphate.

There are other references, mainly of Japanese origin, which claim the use of chondroitin sulphate in preparations for topical application to human skin, particularly as hair tonics.

#### Field of the invention

It is believed that molecules having a molecular weight of greater than about 1500 are unlikely to penetrate skin to any significant extent when applied topically, and accordingly, it is unlikely that polysaccharides such as chondroitin sulphate having a molecular weight of many thousands could penetrate the skin of the scalp, following topical application, and diffuse into the immediate environment of the hair follicle in order to initiate or enhance hair growth. It has however now been discovered that esterified oligosaccharides of smaller molecular weight, including both the novel esterified oligosaccharides according to

the invention, and certain related disaccharides which are known for use in compositions other than for the treatment of baldness, can diffuse through the outer layers of the skin to the immediate environment of the hair follicle,  
5 where they can positively increase hair growth.

It has accordingly been shown that application of these materials to mammalian skin in the region of vellus hair can convert vellus hair to growth as terminal hair.  
10 It has also been shown that the rate of terminal hair growth in mammalian species can be increased by applying these materials to mammalian skin in the region of terminal hair.

15 It has also been shown that the molecular charge and polarity of a selected esterified oligosaccharide will influence its skin penetration. Accordingly, it is apparent that the more polar or highly charged the esterified oligosaccharide, the less likely it is to  
20 penetrate the skin, and for this reason it is advantageous to employ oligosaccharides for this purpose which are esterified to an extent that will reduce their molecular charge. The preferred esterified disaccharides that are selected for optimum penetration of the skin of the scalp  
25 are those which have at least 2, most preferably at least 6 ester groups per molecule.

#### Definition of the invention

30 The invention accordingly also provides compositions suitable for topical application to mammalian skin, particularly to the human scalp, which compositions comprise an oligosaccharide containing hexosamine and uronic acid residues, or mixtures of such oligosaccharides  
35 together with a cosmetically and physiologically acceptable vehicle.

The topical composition

The composition is preferably one comprising one or more esterified oligosaccharides or related disaccharides and a solid, semi-solid or liquid physiologically acceptable vehicle. The nature of the vehicle employed in the preparation of any particular composition will depend on the method intended for administration of that composition. The vehicle can be inert or can possess physiological or pharmaceutical benefits of its own.

The relative amounts of the esterified oligosaccharide of the invention, or related disaccharide, and the vehicle is unimportant, as all that is necessary is the supply of an effective amount of the oligosaccharide or disaccharide for the purpose for which it is administered, the vehicle merely acting as a carrier or diluent of these materials.

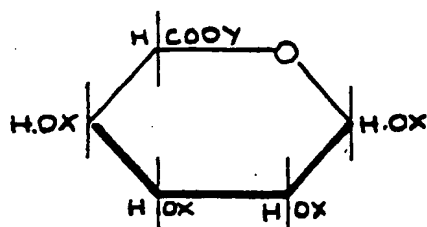
The oligosaccharide

The oligosaccharide that can be incorporated into the composition according to the invention can be one or more of the novel esterified disaccharides as described hereinbefore.

It is also possible to employ esterified oligosaccharides containing three or four or more than four esterified saccharide units, although it is apparent that molecules containing more than four esterified saccharide units may present difficulties, as has been stated earlier, in penetrating the skin to reach the immediate environment of the hair follicle, in view of their excessively large molecular size.

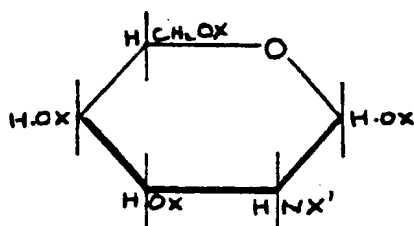
Also suitable for use in the compositions according to the invention are oligosaccharides, especially esterified disaccharides, which are known per se, including those disclosed in EP-A-0 064 012 to Choay S.A., but with a disclosed utility other than for the promotion of hair growth or regrowth.

Examples of such esterified disaccharides described by Choay S.A. are those consisting of a uronic acid residue having the structure:



(50 )

and a hexosamine residue having the structure:



(51 )

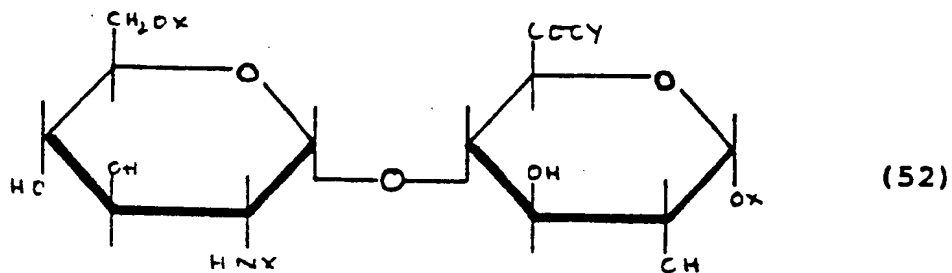
where X is -H, C<sub>1</sub> to C<sub>4</sub> alkyl or an aryl radical,  
 -CO(CH<sub>2</sub>)<sub>m</sub>CH<sub>3</sub>, or -SO<sub>3</sub>M,  
 X' is -H, -CO(CH<sub>2</sub>)<sub>m</sub>CH<sub>3</sub>, or -SO<sub>3</sub>M,  
 Y is -H, C<sub>1</sub> to C<sub>4</sub> alkyl, or M  
 M is -H, or a metallic or organic cation,  
 m is 0 or the integer 1 or 2;



the groups designated X being the same or different, one  
-OH group from each pyranose ring structure being linked  
by a glycosidic linkage having the configuration  $\alpha$ -1,3  
 $\beta$ -1,3  $\alpha$ -1,4 or  $\beta$ -1,4 ; and the -COOY, -CH<sub>2</sub>OX, and -OX  
5 groups being of either configuration with respect to the  
pyranose ring.

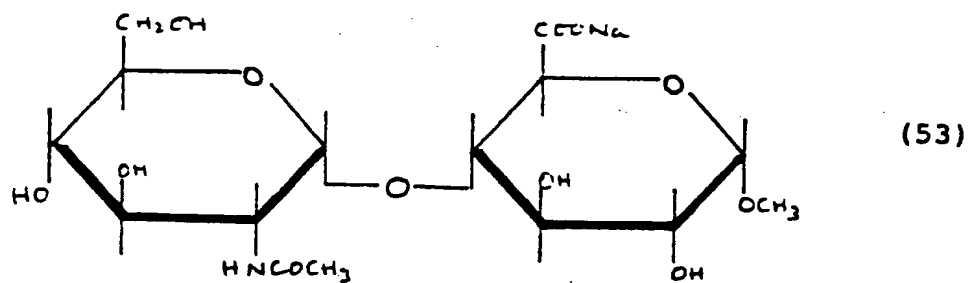
Particularly preferred examples of known esterified  
disaccharides which can be incorporated into compositions  
10 according to the invention have the following structures  
where the designation of X, X', Y and M are as  
hereinbefore defined.

A generic structure of a known esterified disaccharide based on those disclosed in EP-A-0 064 012 to Choay S.A. is:

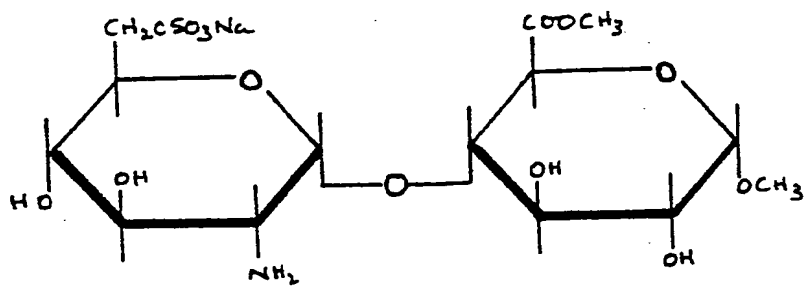


15

Specific examples of the esterified disaccharide (52) have the structures shown as (53) to (59).



5

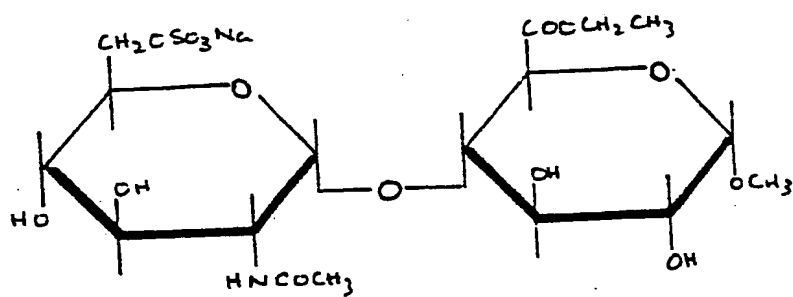


(54)

10

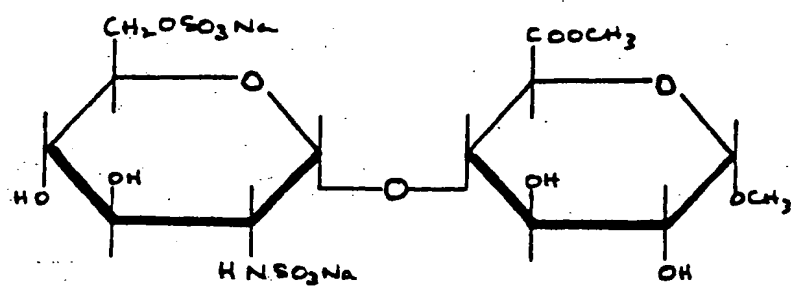
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20



(55)

5

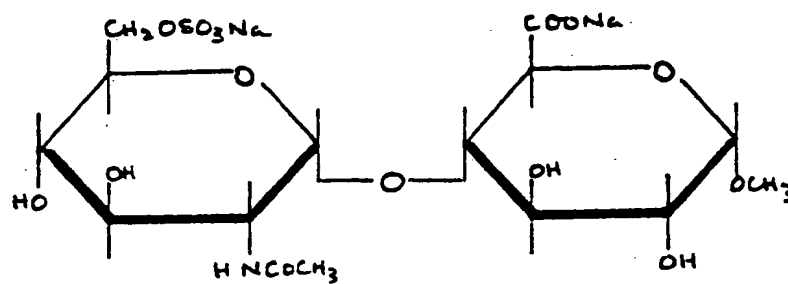


(56)

10

15

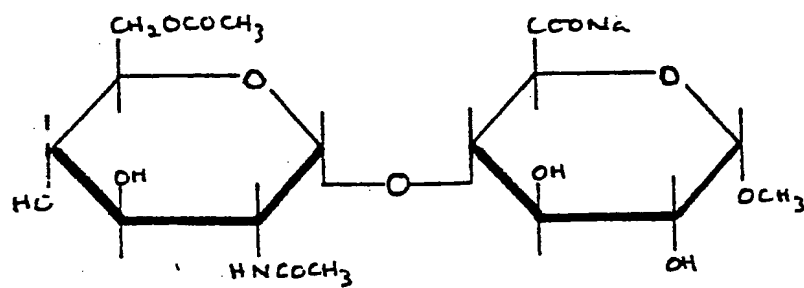
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(57)

5

10

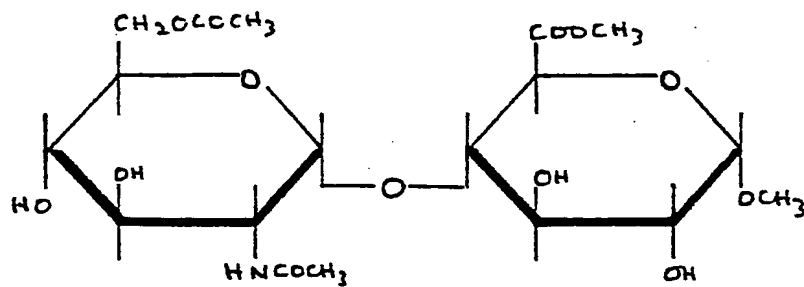


(58)

15

20

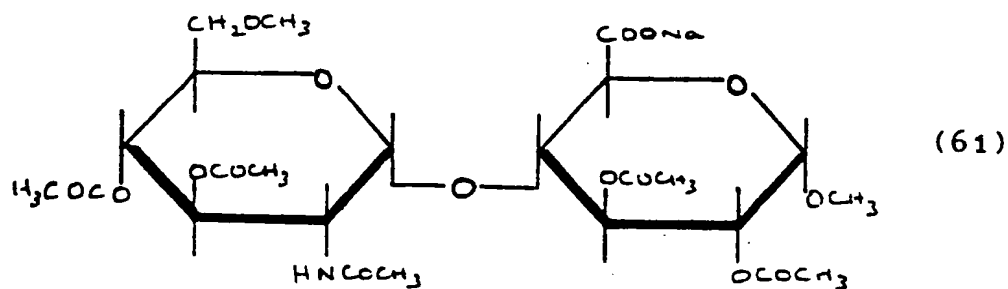
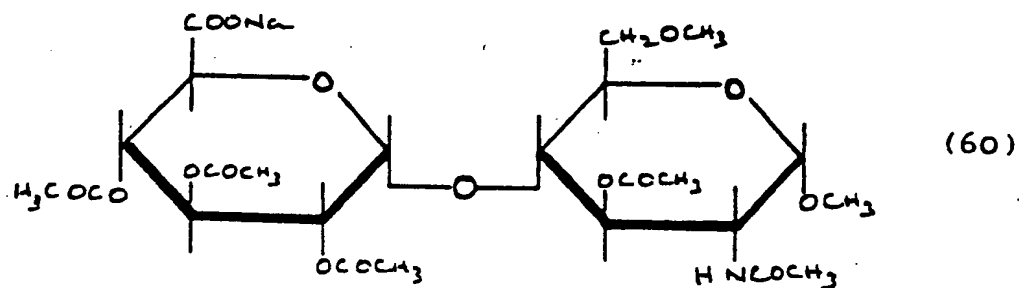
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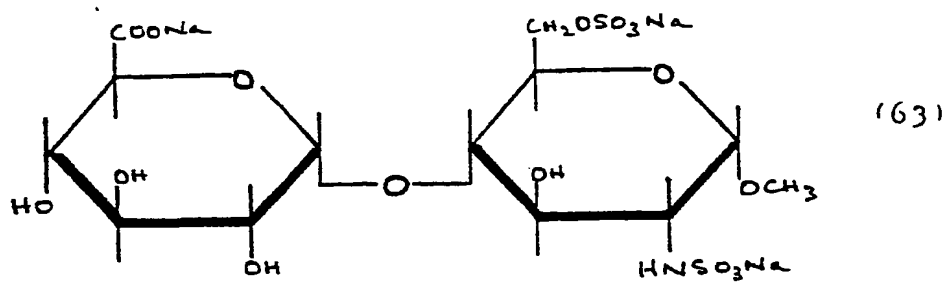
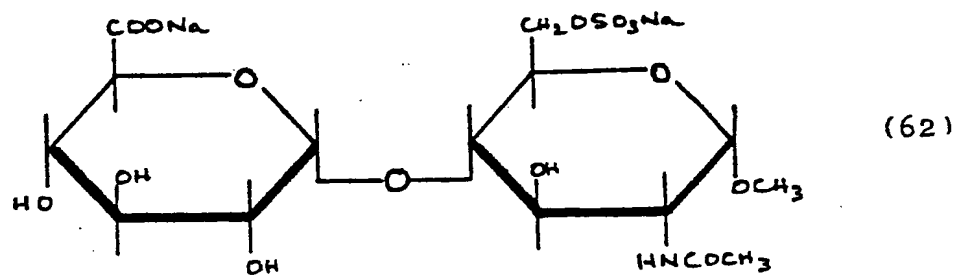


(59)

Specific examples of further esterified disaccharides similar to but not included in the disclosure of EP-A-0 064 012 to Choay S.A., have the structures shown on (60) to (63):

5

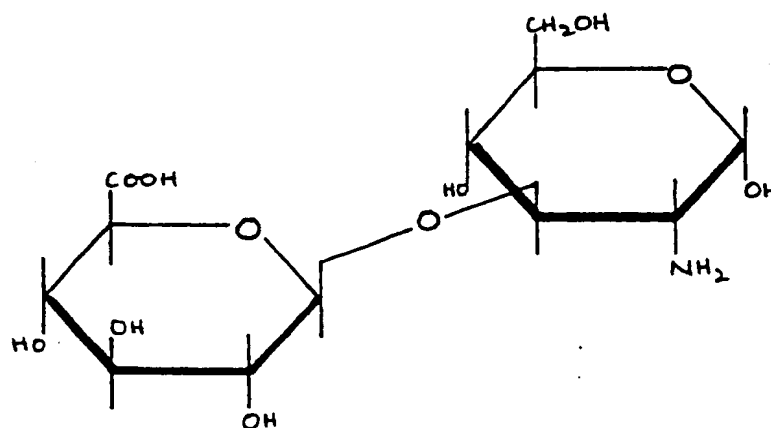




Also suitable for use in compositions according to the invention are oligosaccharides produced by chemical cleavage of glycosaminoglycans, especially chondrosine, having the structure (30), which can be obtained by acid hydrolysis of chondroitin sulphate, and its N-acetyl, and its N-acetyl, O-sulpho derivatives, having the structures (31) and (32) respectively.

Also suitable for use in compositions according to the invention are low molecular weight limit digests of heparan sulphate, hyaluronic acid and chondroitin sulphate produced by the relevant enzyme, namely heparitinase, hyaluronidase and chondroitinase.

Also suitable for use in compositions according to the invention is hyalobiouronic acid having the structure:



(64)



Particularly preferred examples of novel esterified disaccharides according to the invention which can be  
5 incorporated into compositions according to the invention have the structures as shown in hereinafter as (5), (6), (7), (8), (10), (11), (13), (15), (17), (18), (20), (22), (24), (25), (27), (29), (30), (31) and (32).

10 Especially preferred examples of novel esterified oligosaccharides according to the invention, and related disaccharides, which can be incorporated into compositions according to the invention have structures shown hereinbefore as (5), (15), (22) and (31).

15 The amount of the esterified oligosaccharide to be incorporated with a suitable vehicle into compositions for topical use can vary widely, but in general, an amount of from 0.01 to 10%, preferably from 0.1 to 5% by weight of  
20 the composition will provide an adequate dose to the skin after topical application.

#### The Vehicle

25 The composition should also comprise a cosmetically or physiologically acceptable vehicle to enable the esterified oligosaccharide to be conveyed to the skin in an appropriate dilution.

30 The selection of a vehicle for this purpose presents a wide range of possibilities depending on the required product form of the composition. Suitable vehicles can be classified as described hereinafter.

35 It should be explained that vehicles are substances which can act as diluents, dispersants, or solvents for

the esterified oligosaccharide and which therefore ensure that it can be applied to and distributed evenly over the hair and/or scalp at an appropriate concentration. The vehicle is preferably one which can aid penetration of the esterified oligosaccharide into the skin to reach the immediate environment of the hair follicle. Compositions according to this invention can include water as a vehicle, and/or at least one cosmetically acceptable vehicle other than water.

Vehicles other than water that can be used in compositions according to the invention can include solids or liquids such as emollients, solvents, humectants, thickeners and powders. Examples of each of these types of vehicles, which can be used singly or as mixtures of one or more vehicles, are as follows:

Emollients, such as stearyl alcohol, glyceryl monoricinoleate, glyceryl monostearate, propane-1,2-diol, butane-1,3-diol, mink oil, cetyl alcohol, isopropyl isostearate, stearic acid, isobutyl palmitate, isocetyl stearate, oleyl alcohol, isopropyl laurate, hexyl laurate, decyl oleate, octadecan-2-ol, isocetyl alcohol, cetyl palmitate, dimethylpolysiloxane, di-n-butyl sebacate, isopropyl myristate, isopropyl palmitate, isopropyl stearate, butyl stearate, polyethylene glycol, triethylene glycol, lanolin, castor oil, acetylated lanolin alcohols, petroleum, mineral oil, butyl myristate, isostearic acid, palmitic acid, isopropyl linoleate, lauryl lactate, myristyl lactate, decyl oleate, myristylmyristate;

Propellants, such as trichlorofluoromethane, dichlorodifluoromethane, dichlorotetrafluorethane, monochlorodifluoromethane, trichlorotrifluorethane, propane, butane, isobutane, dimethyl ether, carbon dioxide, nitrous oxide;

Solvents, such as ethyl alcohol, methylene chloride, isopropanol, castor oil, ethylene glycol monoethyl ether, diethylene glycol monobutyl ether, diethylene glycol monoethyl ether, dimethyl sulphoxide, tetrahydrofuran;

5

Humectants, such as glycerin, sorbitol, sodium 2-pyrrolidone-5-carboxylate, soluble collagen, dibutyl phthalate, gelatin;

10 Powders, such as chalk, talc, fullers earth, kaolin, starch, gums, colloidal silicon dioxide, sodium polyacrylate, tetra alkyl and/or trialkyl aryl ammonium smectites, chemically modified magnesium aluminium silicate, organically modified montmorillonite clay,  
15 hydrated aluminium silicate, fumed silica, carboxyvinyl polymer, sodium carboxymethyl cellulose, ethylene glycol monostearate.

The amount of vehicle in the composition, including  
20 water if present, should preferably be sufficient to carry at least a portion of the esterified oligosaccharide to the skin in an amount which is sufficient effectively to enhance hair growth. The amount of the vehicle can comprise the balance of the composition, particularly  
25 where little or no other ingredients are present in the composition can accordingly comprise from 10 to 99.99%, preferably from 50 to 99.5% and ideally from 90 to 99% by weight of the vehicle or vehicles.

### 30 Perfume

The composition according to the invention can also optionally comprise a perfume in an amount sufficient to make the composition acceptable to the consumer and  
35 pleasant to use. Usually, the perfume will form from 0.01 to 10% by weight of the composition.

Activity Enhancer

The composition according to the invention can also optionally comprise an activity enhancer whose presence further improves the delivery to the skin of the esterified oligosaccharide. The activity enhancer accordingly effectively increases the partition of the esterified oligosaccharide into the skin from the composition when applied topically.

While screening a series of non-electrolytes for their ability to function as activity enhancers, it was observed that they were all compounds which were capable of substantially increasing the cloud point temperature of nonionic surfactants. Such compounds included short chain alkanols, diols and short chain fatty acids. By contrast, other non-electrolytes which reduce the cloud point temperature of nonionic surfactants, such as long chain alcohols, for example, butan-1-ol and cyclohexanol; polyols, for example sorbitol and propan-1,2,3-triol, do not function as activity enhancers.

An activity enhancer is accordingly defined as a non-electrolyte which is capable, at a concentration of 5% by weight of increasing the cloud point temperature of a 0.025M aqueous solution of polyoxyethylene(8)nonylphenyl ether by at least 10°C. Preferably the activity enhancer is one which is capable of increasing the cloud point temperature by at least 15°C, most preferably by at least 20°C.

It should be explained that the "cloud point" is a measure of the inverse solubility of a nonionic surfactant with temperature and can be determined by heating a standard clear aqueous solution of the nonionic surfactant

until the solution becomes visibly cloudy and then measuring the temperature.

5     The cloud point temperature can conveniently be determined automatically using the equipment and method described by Baum et al in Mat. Res. Std. 4 26 (1964).

10    Examples of suitable compounds functioning as activity enhancers, together with the respective elevated cloud point temperature obtained in each case when using the standard test defined hereinbefore are listed below in Table 1.

Table 1

Elevation of the cloud point temperature of a 0.025M  
aqueous solution of polyoxyethylene(8)nonylphenyl ether  
in the presence of activity enhancers

0.025M aqueous solution of polyoxyethylene(8)nonyl phenyl ether <u>(SYNPERONIC NP8 ex ICI)</u>		
<u>Activity Enhancer</u> <u>(5% by weight)</u>	<u>Cloud point</u> <u>temperature</u> <u>(°C)</u>	<u>Elevation of</u> <u>cloud point</u> <u>temperature</u>
None (control)	33.5	0
2-methyl propan-2-ol	55.0	21.5
Propan-2-ol	56.0	22.5
Ethyl-hydroxypropanoate	52.1	18.6
Hexan-2,5-diol	52.0	18.5
POE(2) ethyl ether	46.0	12.5
Di(2-hydroxypropyl) ether	44.8	11.3
Pentan-2,4-diol	48.0	14.5
Acetone	46.3	12.8
POE(2) methyl ether	43.5	10.0
2-hydroxypropionic acid	45.0	11.5
Propan-1-ol	53.0	19.5
1,4 Dioxane	44.0	10.5
Tetrahydrofuran	45.0	11.5
Butan-1,4-diol	45.0	11.5

Other ingredients

5       The composition according to the invention optionally can contain ingredients other than those already mentioned depending on the form of the intended product. It is, for example, possible to include antiseptics, preservatives, antioxidants, emulsifiers, colouring agents and detergents. It is also possible to include other materials which influence hair regrowth such as minoxidil and derivatives thereof.

10

15       The composition according to the invention can also be employed as a vehicle for a wide variety of cosmetically or pharmaceutically active ingredients, particularly ingredients which have some beneficial effect when applied to the skin other than the promotion of hair growth.

20       The total amount of optional ingredients can form the balance of up to 84.99% by weight of the composition.

Process for preparation of composition

25       The invention also provides a process for the preparation of a composition suitable for topical application to mammalian skin, particularly to the hair and/or scalp of the human subject which comprises mixing an esterified oligosaccharide with a suitable cosmetically or physiologically acceptable vehicle, to provide a composition containing the oligosacchharide at a concentration of from 0.01 to 10% by weight.

30

Form of composition

5 The composition of the invention can be formulated as liquids, for example as a lotion, shampoo, milk or cream for use in conjunction with an applicator such as a roll-ball applicator, or a spray device such as an aerosol can containing propellant, or a container fitted with a pump to dispense the liquid product. Alternatively, the composition of the invention can be solid or semi-solid, 10 for example sticks, creams or gels, for use in conjunction with a suitable applicator or simply a tube, bottle or lidded jar, or as a liquid-impregnated fabric, such as a tissue wipe.

15 The invention accordingly also provides a closed container containing a composition as herein defined.

Use of compositions

20 The invention also provides for the use of an esterified oligosaccharide in the topical treatment of baldness.

25 The compositions according to the invention are primarily intended for topical application to mammalian skin, particularly to the scalp of the human subject, especially where the head is already bald or balding. The compositions can also be applied prophylactically to the hair and hence the scalp to reduce or prevent the 30 onset of baldness.

The amount of the composition and the frequency of application to the hair and/or scalp can vary widely, depending on personal needs, but it is suggested as an 35 example that topical application of from 1 to 5g daily containing from 0.001 to 1g of a selected esterified oligosaccharide over the period of at least six months will in most cases result in an improvement in hair growth.



Efficacy testing

5 The efficacy of the compositions according to the invention was examined using the rat as an animal model, in order to assess both the duration and amount of hair growth, following daily administration of such compositions.

10 For this purpose, hair was clipped from the treatment area at regular intervals and weighed as an index of hair production. The increased duration of anagen phase was apparent from growth curves obtained by plotting weight of hair recovered from clippings against duration of the experiment.

15 This model was validated using minoxidil, a compound known to stimulate hair growth, as a positive control.

20 Effective compounds were selected on the basis of their ability to increase hair production and to increase the duration of anagen phase, both of which are required attributes of any product for reversing baldness.

EXAMPLES

25 The invention is illustrated by the following examples:

Example 1

30 This Example illustrates a lotion according to the invention which is suitable for topical application to the scalp in order to promote hair growth.

The lotion has the following formulation:

	<u>% w/w</u>
disaccharide structure (53)	0.1
ethanol	99.995
perfume	q.s.

5

Example 2

This Example illustrates a hair tonic which is suitable for application to hair or scalp.

10

The hair tonic has the following formulation:

	<u>% w/w</u>
disaccharide : structure (54)	0.8
ethanol	50
water	49
perfume	q.s.

15

Example 3

This Example also illustrates a lotion which is suitable for topical application to the scalp.

20

The lotion has the following formulation:

	<u>% w/w</u>
disaccharide : structure (55)	1.5
propan-2-ol	10
ethanol	88.5
perfume	q.s.

25

Example 4

This Example also illustrates a hair tonic which is suitable for application to hair or scalp.

30

The hair tonic has the following formulation:

	<u>% w/w</u>
disaccharide : structure (56)	0.2
ethanol	40
water	59.80
perfume	q.s.

35

Examples 5 to 8

The following formulations represent lotions which can be used topically in the treatment of bald or balding male or female heads.

		<u>% w/w</u>			
	<u>5</u>	<u>6</u>	<u>7</u>	<u>8</u>	
	Hydroxyethyl cellulose	0.4	-	0.4	-
	Absolute ethanol	25	25	25	25
10	Propane-1,2-diol	-	-	38.4	38.4
	Butane-1,3-diol	38.4	38.8	-	-
	Paramethyl benzoate	0.2	0.2	0.2	0.2
	Disaccharide				
	structure (57)	5	-	-	-
15	Disaccharide				
	structure (58)	-	1	-	-
	Disaccharide				
	structure (59)	-	-	0.8	-
	Disaccharide				
20	structure (60)	-	-	-	0.6
	Perfume	1	1	1	1
	Water	to 100	100	100	100

Examples 9 to 12

25

The following formulations represent creams which can be used in the treatment of baldness.

		<u>8 w/w</u>		
	<u>9</u>	<u>10</u>	<u>11</u>	<u>12</u>
	Cetyl alcohol			
	polyoxyethylene (10)	4	4	4
5	Cetyl alcohol	4	4	4
	Mineral oil	4	2	-
	Paraffin wax	-	2	4
	Partial glyceride of palmitic and			
10	stearic acids	-	-	4
	Disaccharide :			
	structure (30)	2	-	-
	Disaccharide :			
	structure (61)	-	-	1
15	Disaccharide :			
	structure (62)	-	1.5	-
	Disaccharide :			
	structure (63)	-	-	2
	Triethanolamine	0.75	0.75	0.75
20	Butane-1,3-diol	3	3	3
	Xanthan gum	0.3	0.3	0.3
	Preservative	0.4	0.4	0.4
	Perfume	q.s.	q.s.	q.s.
	Water	to 100	100	100
25				

### Example 13

30 This Example illustrates a water-in-oil high internal phase emulsion containing a novel disaccharide according to the invention.

The emulsion consisted of 10% by volume oily phase and 90% by weight aqueous phase.

35 The oily phase and the aqueous phase had the following constitution:

	<u>% w/w</u>
<u>Oily phase</u>	
Sorbitan monooleate	20
5 Quartenium-18 hectorite	5
Liquid paraffin	75
<u>Aqueous phase</u>	
Disaccharide : structure (6)	0.5
10 Xanthan gum	1
Preservative	0.3
Perfume	q.s.
Sodium chloride (1% w/w solution)	to 100

15        The emulsion was prepared by taking 10 parts by volume of the oily phase and to it adding slowly with stirring 90 parts by volume of the aqueous phase.

20        The high internal phase water-in-oil emulsion so formed can be applied topically to the scalp, to improve hair growth and regrowth.

25        The following examples 14 to 18 illustrate shampoos for use in washing the hair and scalp, and for promoting hair growth on the scalp.

Example 14

	<u>% w/w</u>
Sodium lauryl ether sulphate	
(2 EO) : 21% AD	41.4
5 Lauryl dimethylamino acetic acid	
betaine * 30% AD	4
Coconut fatty acid diethanolamine	1.5
Oleyl triethoxy phosphate (BRIPHOS 03D)	1
Polyglycol-polyamine condensation	
10 resin (POLYQUART H) : 50% active	1.5
Preservative, colouring matter, salt	0.58
Disaccharide : structure (7)	5
Perfume	q.s.
Water	to 100

Example 15

	<u>% w/w</u>
Sodium lauryl ether sulphate (2 EO) :	
100% AD	12
POLYQUART H : 50% active	2.5
20 BRIPHOS 03D	2.5
Disaccharide : structure (8)	4
Perfume	q.s.
Water	to 100

Example 16

	<u>% w/w</u>
25 Monoethanolamine lauryl sulphate :	
100% AD	20
POLYQUART H : 50% active	3
BRIPHOS 03D	1.7
30 Coconut diethanolamide	5
Disaccharide : structure (5)	1
Perfume	q.s.
Water	to 100
pH adjusted to 6.5	

Example 17

	<u>% w/w</u>
Sodium lauryl ether sulphate (3 EO) :	
5     100% AD	12
POLYQUART H : 50% active	0.3
ERIPHOS 03D	1
Disaccharide : structure (10)	2
Perfume	q.s.
10    Water	to 100

pH adjusted to 6.5

Example 18

	<u>% w/w</u>
Sodium lauryl ether sulphate ( 2 EO) :	
100% AD	12
POLYQUART H : 50% active	3
20    ERIPHOS 03D	1
Cpacifier	9
Disaccharide : structure (11)	5
Perfume	q.s.
Water	to 100

25

pH adjusted to 6.5

Examples 19 to 24

30     The following Examples 19 to 24 illustrate powder compositions according to the invention which can be applied topically to the scalp.

		<u>19</u>	<u>20</u>	<u>21</u>	<u>% w/w</u>		<u>22</u>	<u>23</u>	<u>24</u>
	Chemically modified								
	starch	5	-	5	-		5	-	
5	Chemically modified								
	cellulose	-	5	-	5	-		5	
	Boric acid	10	10	10	10	10	10	10	
	Zinc oxide	5	5	5	5	5	5	5	
	Disaccharide :								
10	structure (13)	5	-	-	-	-	-	-	
	Disaccharide :								
	structure (15)	-	10	-	-	-	-	-	
	Disaccharide :								
	structure (18)	-	-	2	-	-	-	-	
15	Disaccharide :								
	structure (17)	-	-	-	4	-	-	-	
	Disaccharide :								
	structure (20)	-	-	-	-	1	-	-	
	Disaccharide :								
20	structure (22)	-	-	-	-	-	-	3	
	Perfume	q.s.	q.s.	q.s.	q.s.	q.s.	q.s.	q.s.	
	Chalk	10	10	10	10	10	10	10	
	Talc	to 100	100	100	100	100	100	100	

Example 25

25

This Example illustrates a lotion according to the invention which is suitable for topical application to the scalp in order to promote hair growth.

30

The lotion has the following formulation:

		<u>% w/w</u>
	disaccharide structure (24)	0.1
	ethanol	99.995
35	perfume	q.s.



Example 26

This Example illustrates a hair tonic which is suitable for application to hair or scalp.

5

The hair tonic has the following formulation:

	<u>% w/w</u>
disaccharide : structure (25)	0.9
10 ethanol	50
water	49
perfume	q.s.

Example 27

15

This Example also illustrates a lotion which is suitable for topical application to the scalp.

The lotion has the following formulation:

	<u>% w/w</u>
disaccharide : structure (27)	1.5
propan-2-ol	10
ethanol	88.5
perfume	q.s.

25

Example 28

This Example also illustrates a hair tonic which is suitable for application to hair or scalp.

30

The hair tonic has the following formulation:

	disaccharide : structure (29)	<u>% w/w</u>
	ethanol	0.2
	water	40
5	perfume	59.80
		q.s.

Examples 29 to 32

The following formulations represent lotions which can be used topically in the treatment of bald or balding male or female heads.

		<u>% w/w</u>			
		<u>29</u>	<u>30</u>	<u>31</u>	<u>32</u>
	Hydroxyethyl cellulose	0.4	-	0.4	-
15	Absolute ethanol	25	25	25	25
	Propane-1,2-diol	-	-	38.4	38.4
	Butane-1,3-diol	38.4	38.8	-	-
	Paramethyl benzoate	0.2	0.2	0.2	0.2
	Disaccharide				
20	structure (30)	5	1	-	-
	Disaccharide				
	structure (31)	-	1	4	-
	Disaccharide				
	structure (32)	-	-	1	3
25	Disaccharide				
	structure (33)	2	-	-	3
	Perfume	1	1	1	1
	Water	to 100	100	100	100

30

Examples 33 to 36

The following formulations represent creams which can be used in the treatment of baldness.

		<u>% w/w</u>			
		<u>33</u>	<u>34</u>	<u>35</u>	<u>36</u>
	Cetyl alcohol				
	polyoxyethylene (10)	4	4	4	4
5	Cetyl alcohol	4	4	4	4
	Mineral oil	4	2	-	-
	Paraffin wax	-	2	4	-
	Partial glyceride				
	of palmitic and				
10	stearic acids	-	-	-	4
	Disaccharide :				
	structure (10)	4	-	2	-
	Disaccharide :				
	structure (11)	-	5	-	1
15	Disaccharide :				
	structure (13)	-	1.5	-	3.5
	Disaccharide :				
	structure (18)	2	-	2	-
	Triethanolamine	0.75	0.75	0.75	0.75
20	Butane-1,3-diol	3	3	3	3
	Xanthan gum	0.3	0.3	0.3	0.3
	Preservative	0.4	0.4	0.4	0.4
	Perfume	q.s.	q.s.	q.s.	q.s.
	Water	to 100	100	100	100

25

Example 37

30 This Example illustrates a water-in-oil high internal phase emulsion containing a novel disaccharide according to the invention.

The emulsion consisted of 10% by volume oily phase and 90% by weight aqueous phase.

35 The oily phase and the aqueous phase had the following constitution:

		<u>% w/w</u>
	<u>Oily phase</u>	
	Sorbitan monooleate	20
	Quartenium-18 hectorite	5
5	Liquid paraffin	75

	<u>Aqueous phase</u>	
	Disaccharide : structure (36)	0.5
	Xanthan gum	1
10	Preservative	0.3
	Perfume	q.s.
	Sodium chloride (1% w/w solution)	to 100

15 The emulsion was prepared by taking 10 parts by volume of the oily phase and to it adding slowly with stirring 90 parts by volume of the aqueous phase.

20 The high internal phase water-in-oil emulsion so formed can be applied topically to the scalp, to improve hair growth and regrowth.

25 The following examples 38 to 42 illustrate shampoos for use in washing the hair and scalp, and for promoting hair growth on the scalp.

Example 38

		<u>% w/w</u>
	Sodium lauryl ether sulphate	
5	(2 EO) : 21% AD	41.4
	Lauryl dimethylamino acetic acid	
	betaine: 30% AD	4
	Coconut fatty acid diethanolamine	1.5
	Oleyl triethoxy phosphate (BRIPHOS 03D)	1
10	Polyglycol-polyamine condensation	
	resin (POLYQUART H) : 50% active	1.5
	Preservative, colouring matter, salt	0.58
	Disaccharide : structure (37)	5
	Perfume	q.s.
15	Water	to 100

Example 39

		<u>% w/w</u>
20	Sodium lauryl ether sulphate (2 EO) :	
	100% AD	12
	POLYQUART H : 50% active	2.5
	BRIPHOS 03D	2.5
	Disaccharide : structure (15)	4
25	Perfume	q.s.
	Water	to 100

Example 40

	<u>% w/w</u>
Monoethanolamine lauryl sulphate :	
100% AD	20
5 POLYQUART H : 50% active	3
BRIPHOS 03D	1.7
Coconut diethanolamide	5
Disaccharide : structure (56)	1
Perfume	q.s.
10 Water	to 100
pH adjusted to 6.5	

Example 41

	<u>% w/w</u>
15 Sodium lauryl ether sulphate (3 EO) :	
100% AD	12
POLYQUART H : 50% active	0.3
BRIPHOS 03D	1
Disaccharide : structure (57)	2
20 Perfume	q.s.
Water	to 100
pH adjusted to 6.5	

Example 42

	<u>% w/w</u>
Sodium lauryl ether sulphate ( 2 EO ) :	
100% AD	12
5   POLYQUART H : 50% active	3
BRIPHOS 03D	1
Opacifier	9
Disaccharide : structure (58)	5
Perfume	q.s.
10   Water	to 100
pH adjusted to 6.5	

Examples 43 to 47

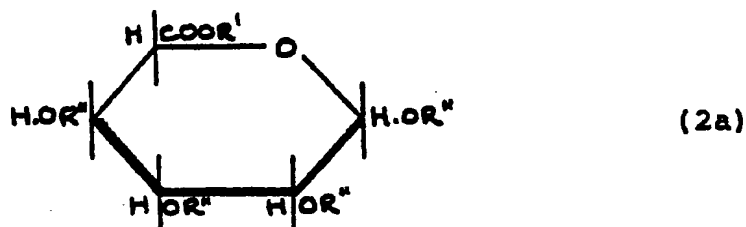
The following Examples 43 to 47 illustrate powder compositions according to the invention which can be applied topically to the scalp.

		<u>% w/w</u>				
		<u>43</u>	<u>44</u>	<u>45</u>	<u>46</u>	<u>47</u>
5	Chemically modified starch	5	-	5	-	5
	10 Chemically modified cellulose	-	5	-	5	-
	Boric acid	10	10	10	10	10
	Zinc oxide	5	5	5	5	5
	Disaccharide :					
15	structure (59)	5	-	-	-	-
	Disaccharide :					
	structure (60)	-	10	-	-	-
	Disaccharide :					
	structure (61)	-	-	2	-	-
20	Disaccharide :					
	structure (62)	-	-	-	4	-
	Disaccharide :					
	structure (63)	-	-	-	-	1
	Perfume	q.s.	q.s.	q.s.	q.s.	q.s.
25	Chalk	10	10	10	10	10
	Talc	to 100	100	100	100	100



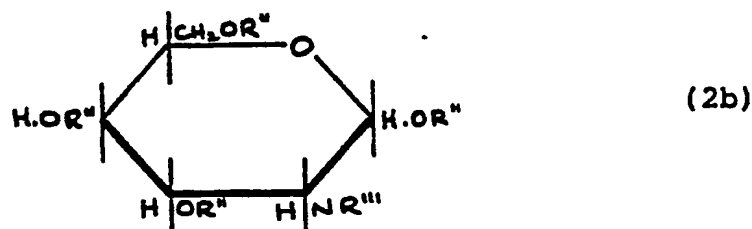
CLAIMS

1. An esterified oligosaccharide including at least one esterified disaccharide unit consisting of a uronic acid residue having the structure:



15

and a hexosamine residue having the structure:

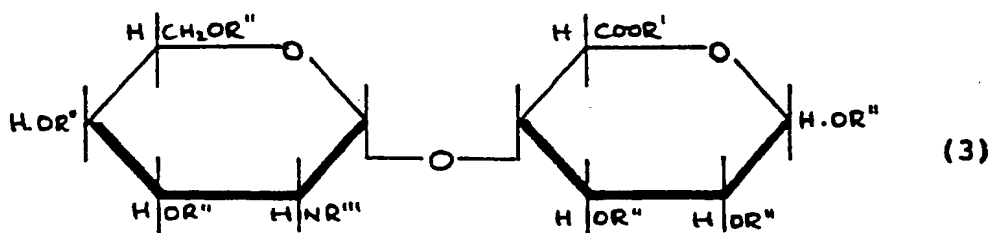


25

Where R' is C<sub>3</sub> to C<sub>10</sub> alkyl or  $\begin{matrix} \text{COOR''} \\ | \\ -\text{CH}(\text{CH}_2)_n \text{CH}_3 \end{matrix}$   
R'' is -H, C<sub>1</sub> to C<sub>4</sub> alkyl,  $-\text{CO}(\text{CH}_2)_m \text{CH}_3$ ,  $-\text{SO}_3\text{M}$   
R''' is -H,  $-\text{CO}(\text{CH}_2)_m \text{CH}_3$ , or  $-\text{SO}_3\text{M}$   
M is -H, or a metallic or organic cation  
n is 0 or an integer of from 1 to 7  
m is 0 or the integer 1 or 2

30

the groups designated R" being the same or different, one R" group from each pyranose ring structure being linked by a glycosidic linkage having the configuration  $\alpha$ -1,3;  $\alpha$ -1,4;  $\beta$ -1,3; or  $\beta$ -1,4; and the -COOR', -CH<sub>2</sub>OR" and -OR" groups being of either configuration with respect to the pyranose rings; provided the when the esterified oligosaccharide is an esterified disaccharide unit having the structure:

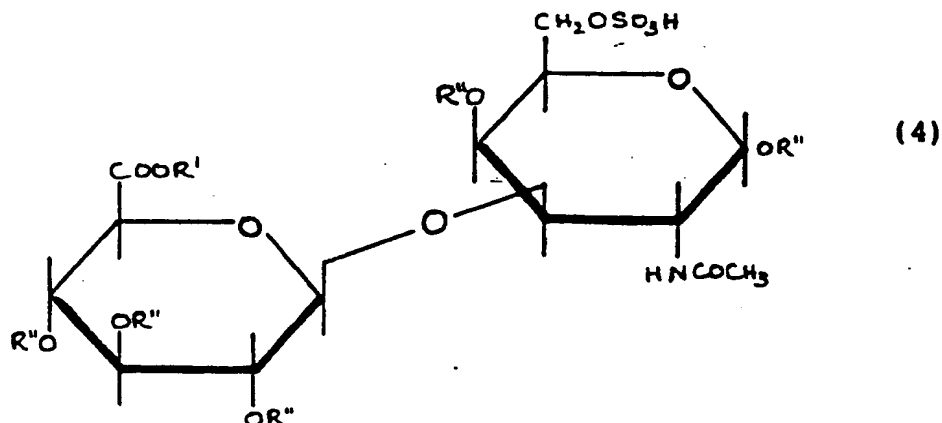


then R' is C<sub>5</sub> to C<sub>10</sub> alkyl or  $\text{-}\overset{\text{COOR}''}{\underset{|}{\text{CH}}}(\text{CH}_2)_n\text{CH}_3$ .

2. An esterified oligosaccharide according to Claim 1 which comprises from 2 to 8 saccharide residues, at least two of which form the esterified disaccharide unit.

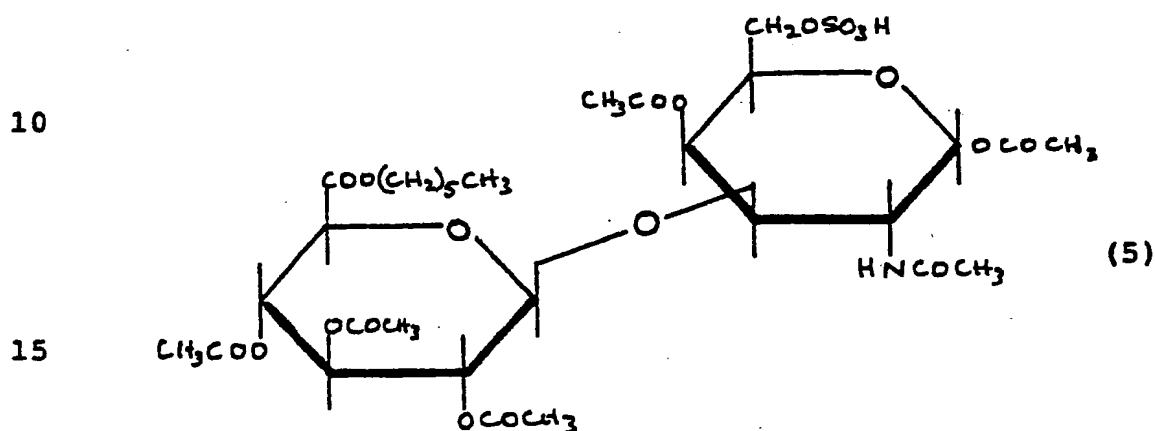
3. An esterified oligosaccharide according to Claim 2, which comprises from 2 to 4 saccharide residues.

4. An esterified oligosaccharide according to any preceding claim having the generic structure:

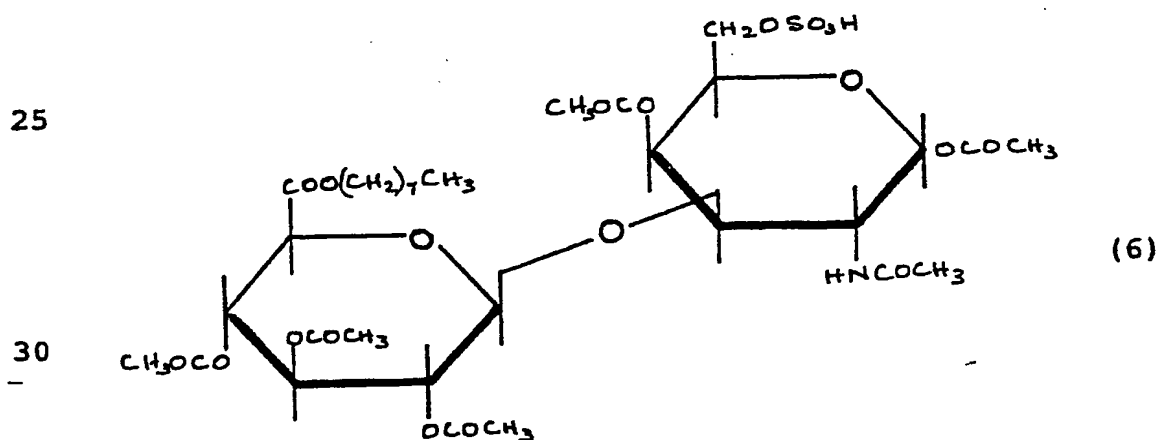


5. An esterified oligosaccharide according to claim 4,  
chosen from those having the following structures:

5



20

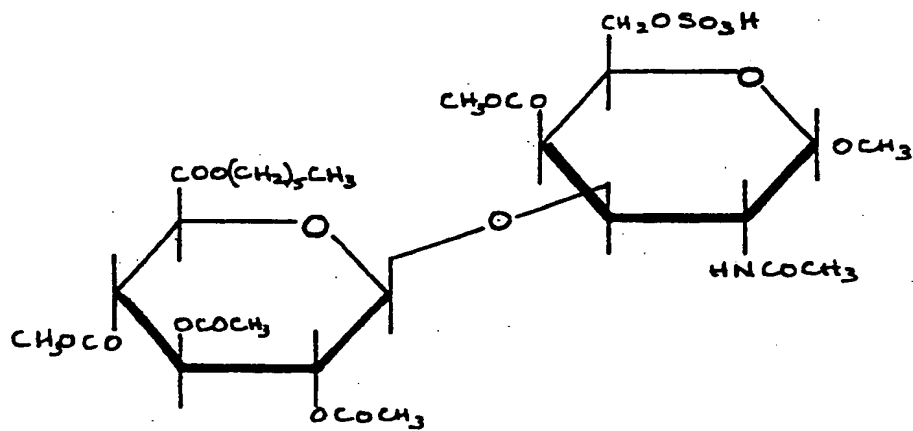


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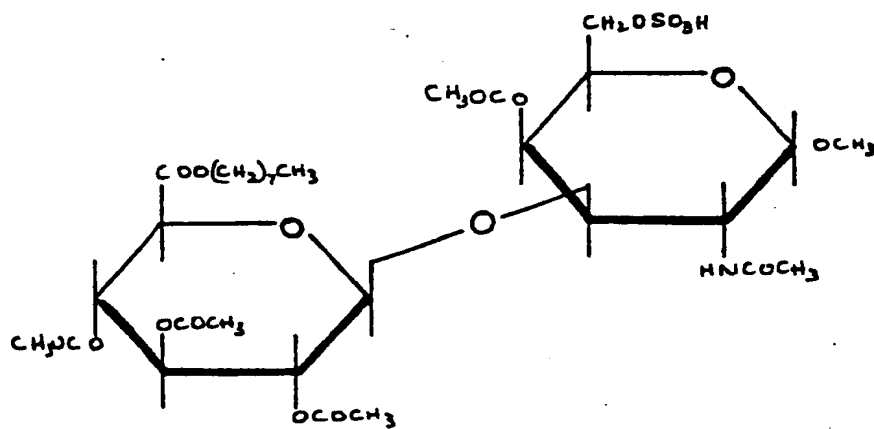


(7)

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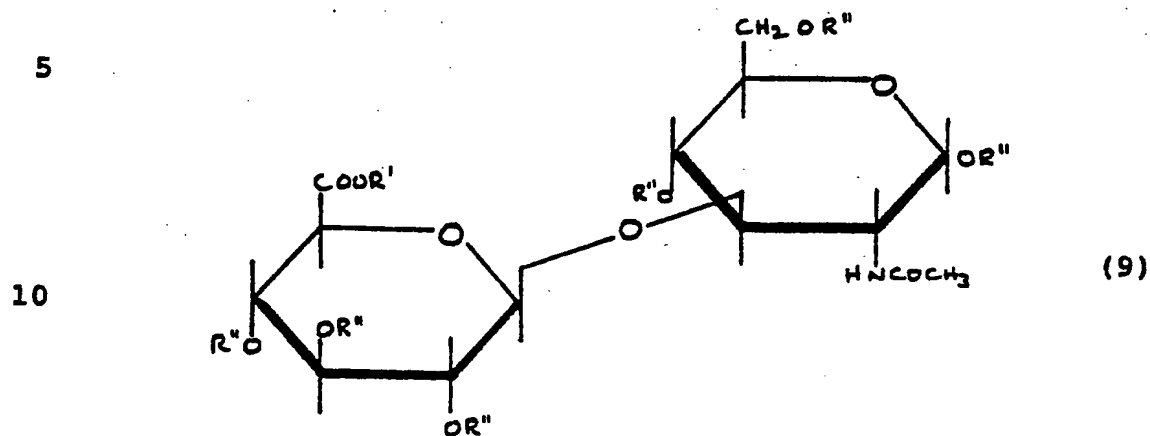
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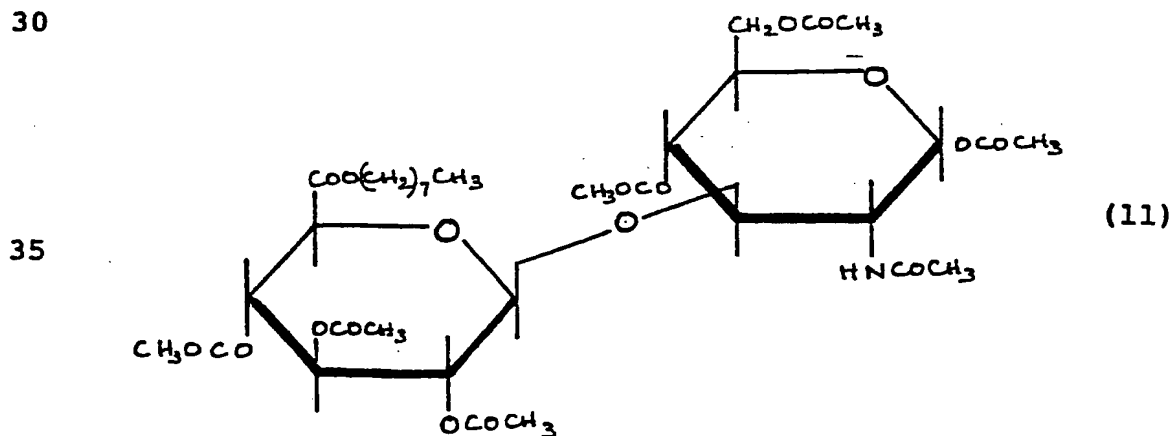
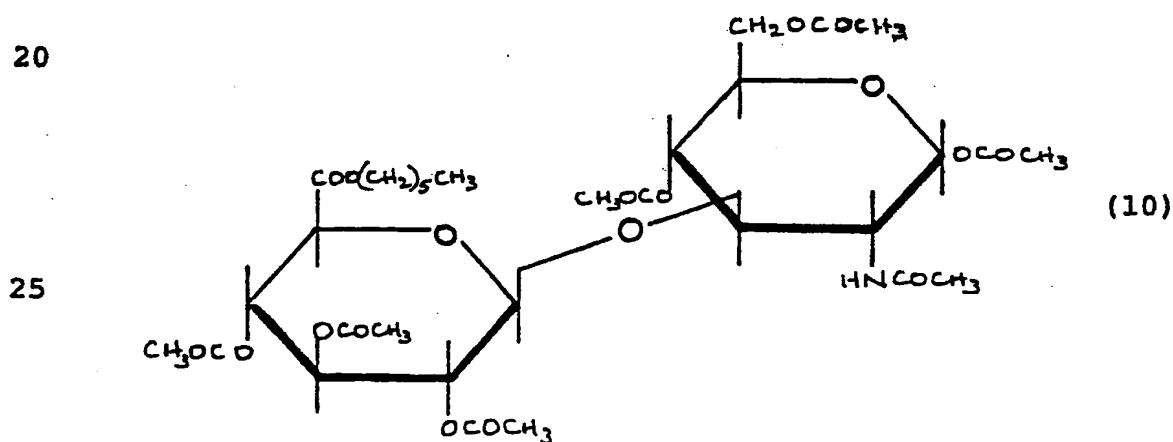


(8)

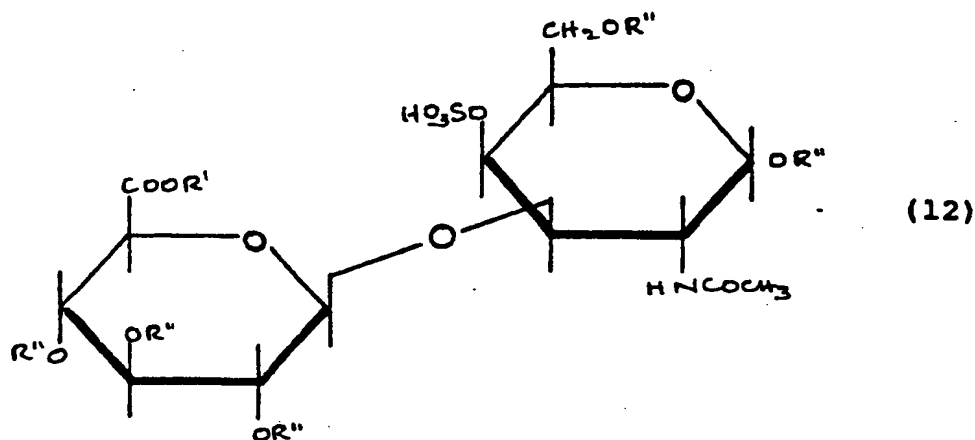
6. An esterified oligosaccharide according to any of claims 1 to 3 having the generic structure:



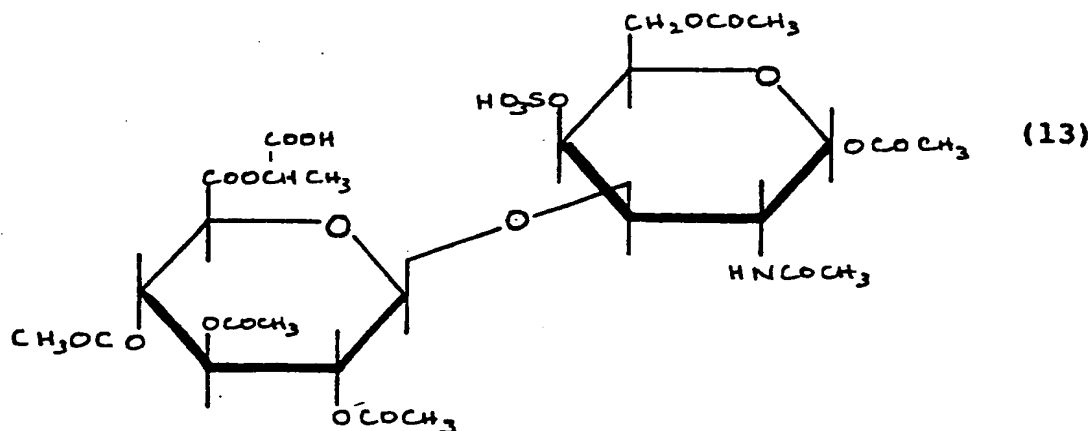
15 7. An esterified oligosaccharide according to Claim 6, chosen from those having the structures:



8. An esterified oligosaccharide according to any of Claims 1 to 3, having the generic structure:



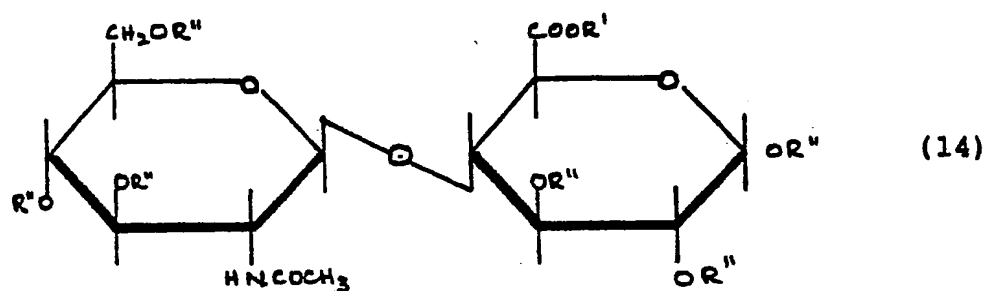
9. An esterified oligosaccharide according to Claim 8 having the structure:



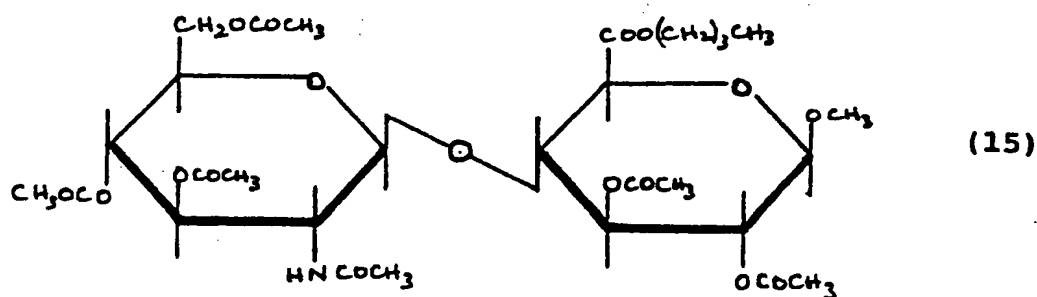
10. An esterified oligosaccharide according to any of Claims 1 to 3, having the generic structure:

5

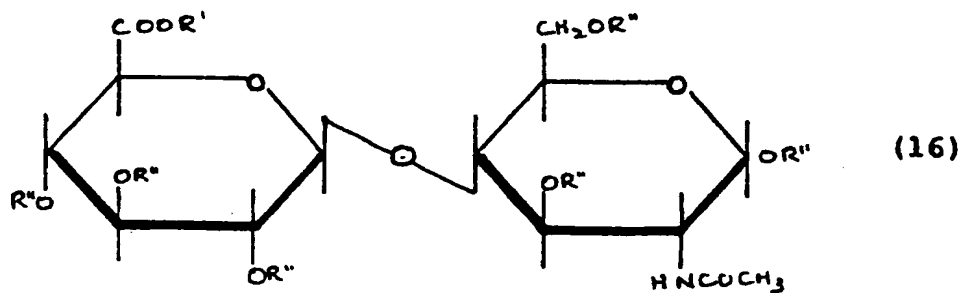
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11. An esterified oligosaccharide according to Claim 10 having the structure:



12. An esterified oligosaccharide according to Claim 1, 2 or 3, having the generic structure:

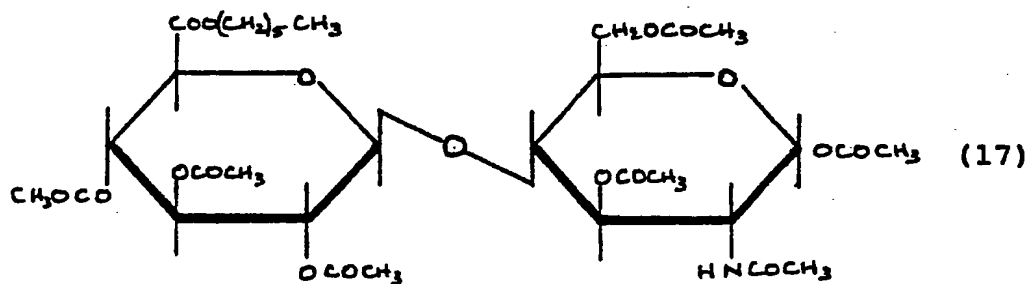




13. An esterified oligosaccharide according to Claim 12  
chosen from those having the structure:

5

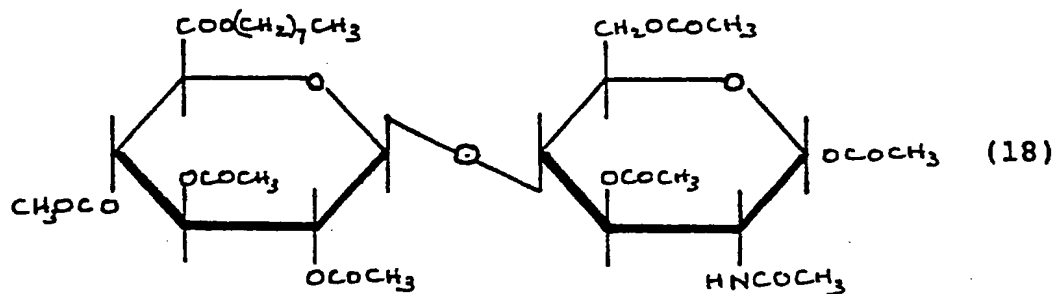
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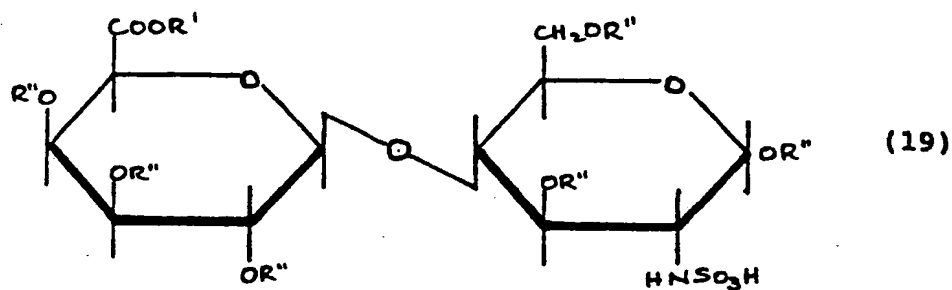
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14. An esterified oligosaccharide according to Claim 1, 2 or 3, having the generic structure:

5

10

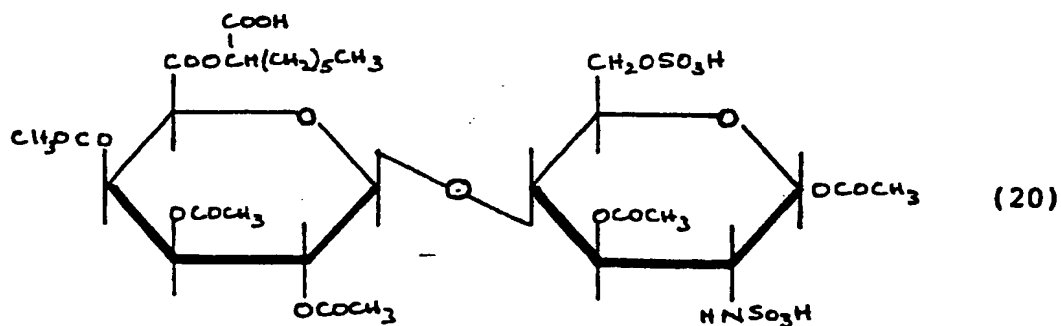


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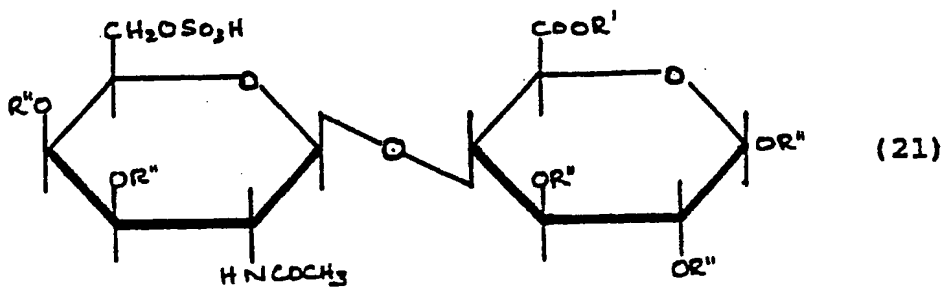
15. An esterified oligosaccharide according to Claim 14  
20 having the structure:

25

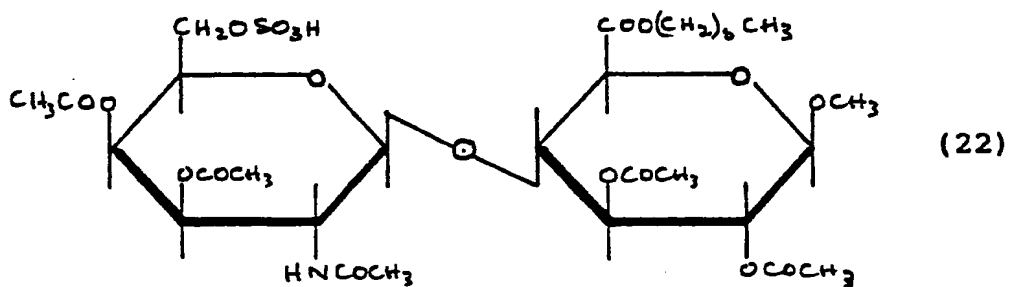
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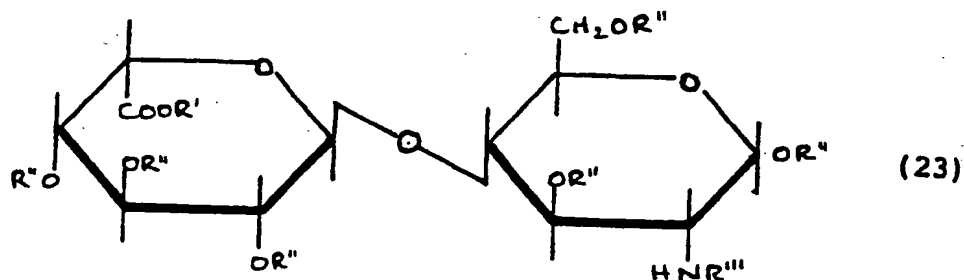
16. An esterified oligosaccharide according to Claim 1, 2 or 3 having the generic structure:



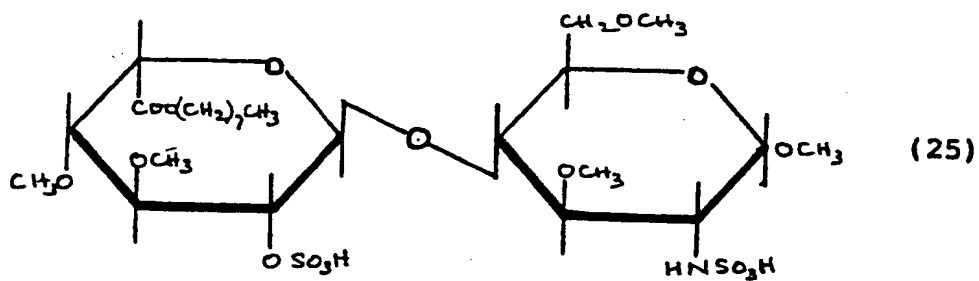
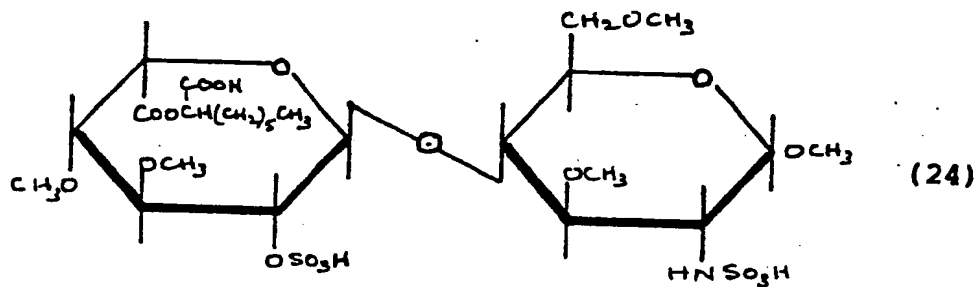
17. An esterified oligosaccharide according to Claim 16, having the structure:



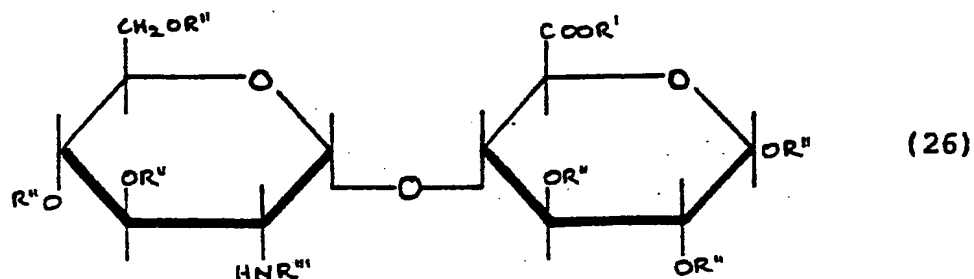
18. An esterified oligosaccharide according to Claim 1, 2 or 3, having the generic structure:



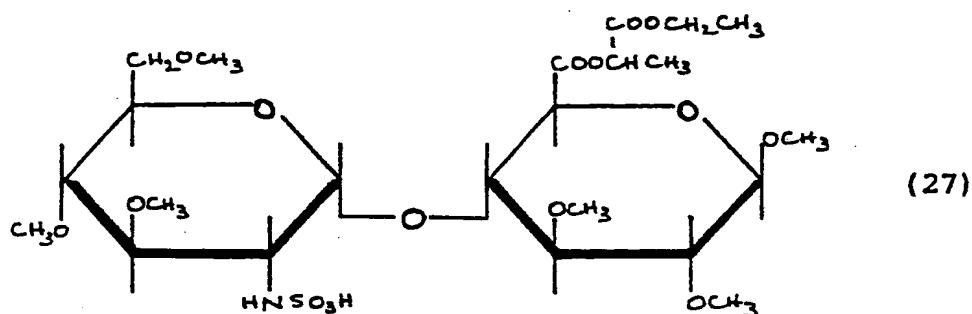
15 19. An esterified oligosaccharide according to Claim 18, chosen from those having the structure:



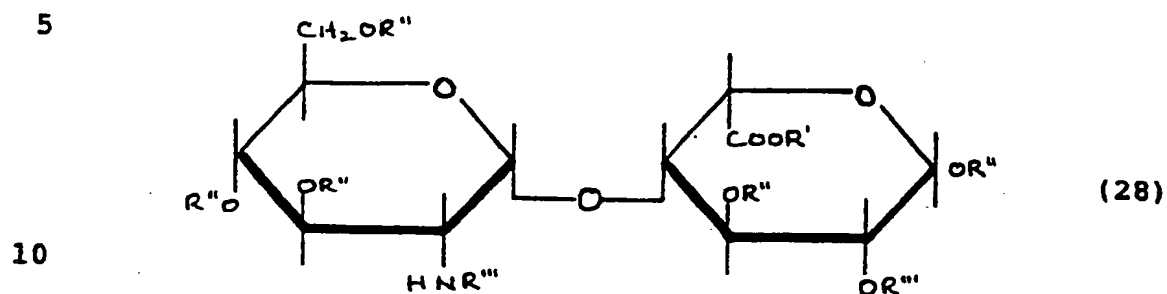
20. An esterified oligosaccharide according to Claim 1, 2 or 3, having the generic structure:



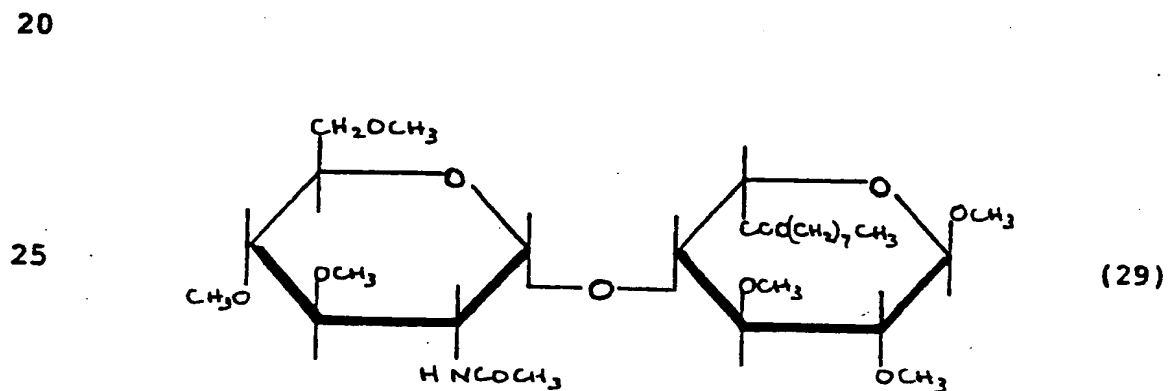
21. An esterified oligosaccharide according to Claim 20, having the structure:



22. An esterified oligosaccharide according to Claim 1, 2 or 3, having the generic structure:



15 23. An esterified oligosaccharide according to Claim 22, having the structure:

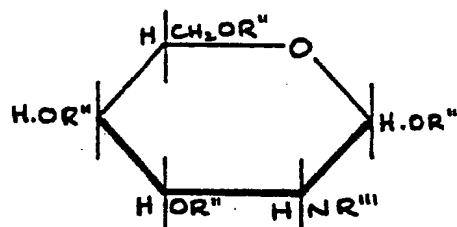


24. A process for the synthesis of an esterified oligosaccharide as defined in any claims 1 to 9, which comprises the steps of:

- 5 ( i) Subjecting a glycosaminoglycan chain to chemical cleavage to yield oligosaccharide fragments containing at least one disaccharide unit comprising a hexosamine residue having the structure:

10

15



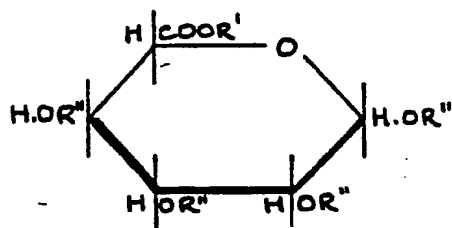
(2b)

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which is glycosidically linked to the C-1 position of a uronic acid residue having the structure:

25

30



(2a)

and (ii) modifying the oligosaccharide fragments by one or more process steps in any sequence chosen from:

acylation of free hydroxyl groups,  
acylation of free amino groups,  
sulphation of free hydroxyl groups,  
sulphation of free amino groups,  
esterification of free hydroxyl groups, and  
etherification of free hydroxyl groups;

to provide the esterified oligosaccharide.

25. A process according to claim 24 for the synthesis of an esterified oligosaccharide as defined in any of claims 1 to 9, having the structure chosen from structures (4), (9) and (12), in which the glycosaminoglycan is chosen from chondroitin sulphate and hyaluronic acid.

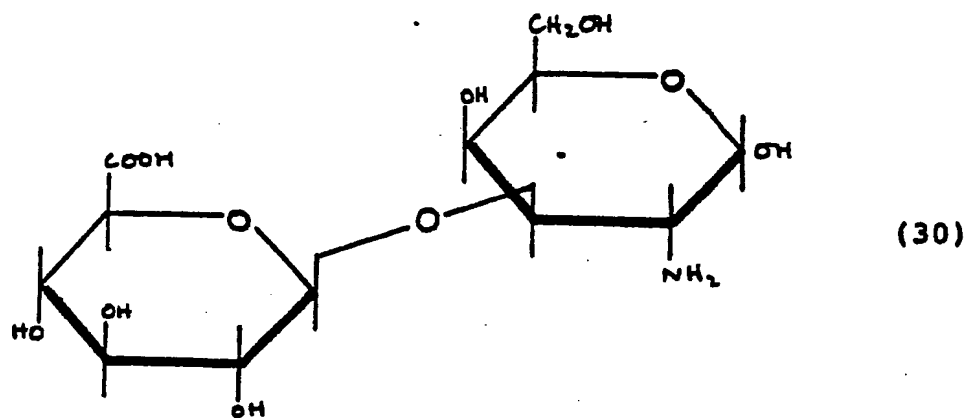
26. A process according to claim 24 or 25, in which the chemical cleavage is achieved by acid hydrolysis.

27. A process according to claim 24 or 25, in which the chemical cleavage is achieved by enzymic digestion.

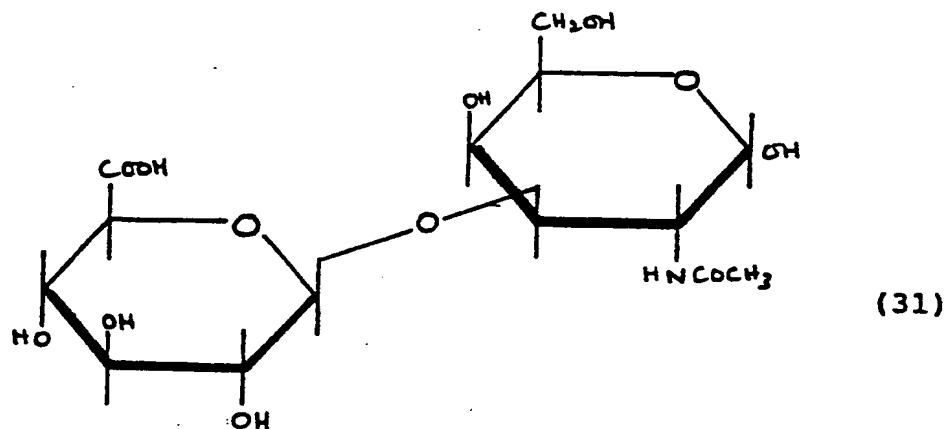


28. A process according to claim 24, 25 or 26, for the synthesis of the esterified disaccharide having the structure (5), which process comprises the steps of:

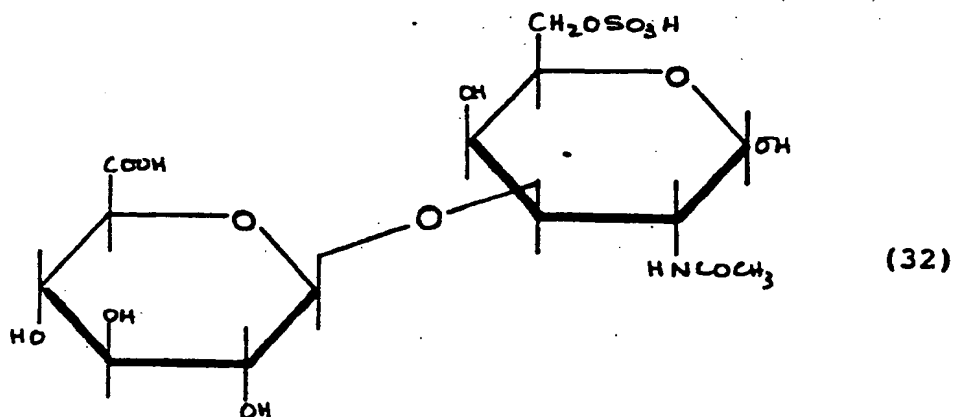
- ( i ) subjecting the glycosaminoglycan chondroitin sulphate to acid hydrolysis to yield the disaccharide chondrosine having the structure;



- ( ii ) subjecting the chondrosine to N-acetylation to yield N-acetylchondrosine having the structure;

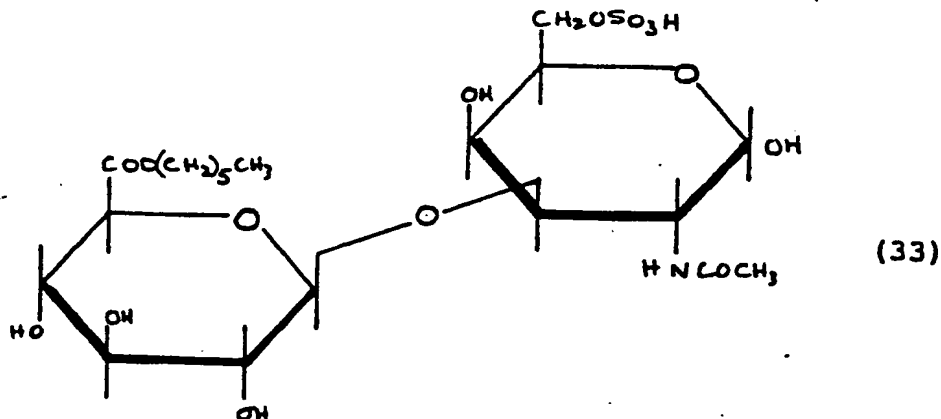


(iii) subjecting the N-acetylchondrosine to  
sulphation in the C-6 position of the hexosamine moiety to  
yield N-acetylchondrosine -6-O-sulphate having the  
5 structure;



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( iv) subjecting the N-acetylchondrosine  
-6-O-sulphate to esterification to yield the intermediate  
product having the structure; and

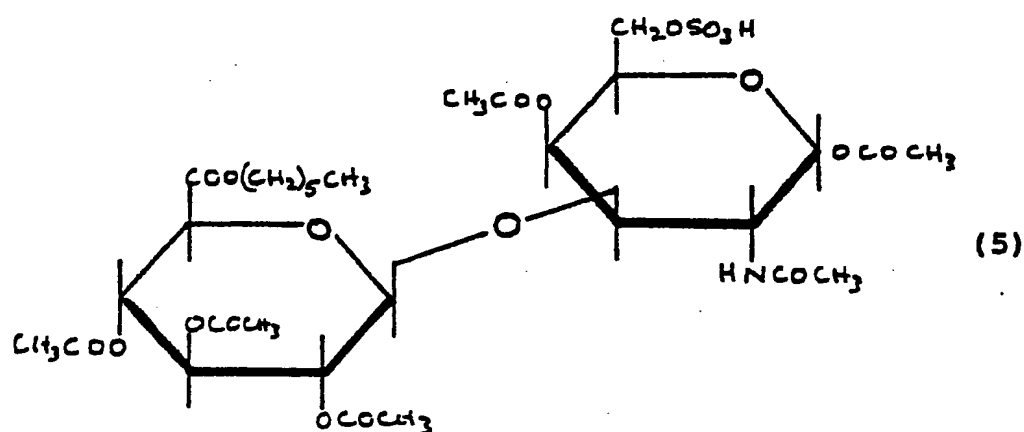


( v) subjecting the intermediate product having the structure (33) to full acetylation to yield the esterified disaccharide having the structure.

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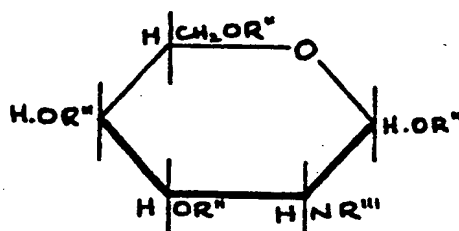
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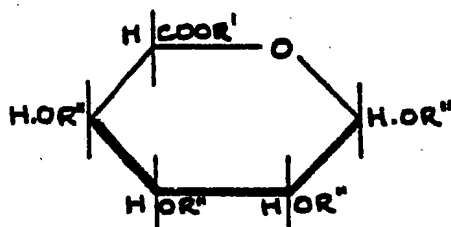
29. A process for the synthesis of an esterified oligosaccharide as defined in any of claims 1 to 3 and 10 to 23 having at least one uronic acid residue linked in the  $\beta$  configuration through the C-1 position to a hexosamine residue, or having at least one hexosamine residue linked in the  $\alpha$  or  $\beta$  configuration through the C-1 position to a uronic acid residue, which process comprises the steps of:

( i) condensing a hexosamine residue having the structure:



(2b)

with a uronic acid residue having the structure:



(2a)

to form an oligosaccharide intermediate, the functional groups of which residues have been modified or protected in such a way that glycosidic bond formation can only occur between the C-1 position of one residue and a free hydroxyl group in the C-3 or C-4 position of the other residue, the  $\alpha$  or  $\beta$  configuration of the glycosidic linkage so formed being dictated by the nature of the leaving group on the C-1 position; and

( ii) removal of at least one protecting group from the oligosaccharide intermediate by one or more process steps in any sequence chosen from:

reduction,  
acid catalysis, and  
base catalysis;

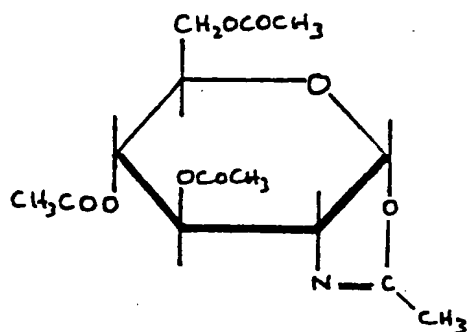
to provide the esterified oligosaccharide.

30. A process according to claim 29, which further comprises modifying the oligosaccharide intermediate by one or more process steps in any sequence, chosen from:

- acylation of free hydroxyl groups
- acylation of free amino groups
- sulphation of free hydroxyl groups
- sulphation of free amino groups
- esterification of free hydroxyl groups, and
- etherification of free hydroxyl groups.

31. A process according to claim 29 or 30, for the synthesis of the esterified disaccharide having the structure (15), which process comprises the steps of:

- ( i ) condensing the oxazoline derivative of N-acetylglucosamine having the structure:

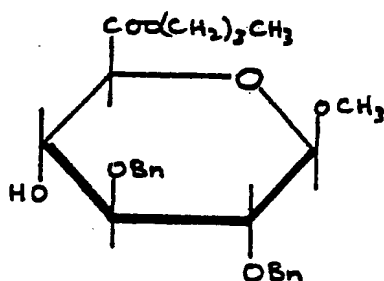


(34)

with the benzylated uronic acid moiety having the structure:

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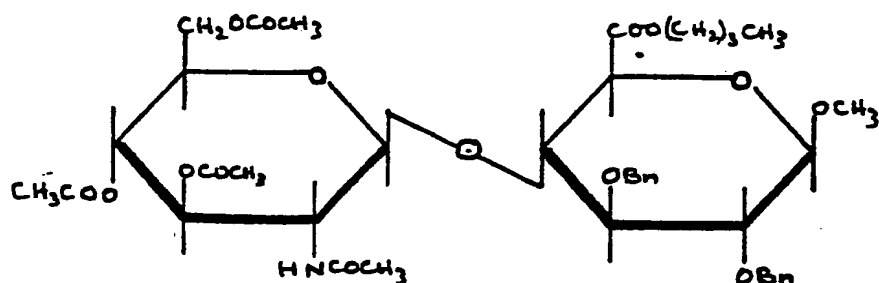
(35)

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in the presence of nitromethane and p-toluene sulphonic acid to yield the benzylated  $\beta$ -1,4 disaccharide having the structure:

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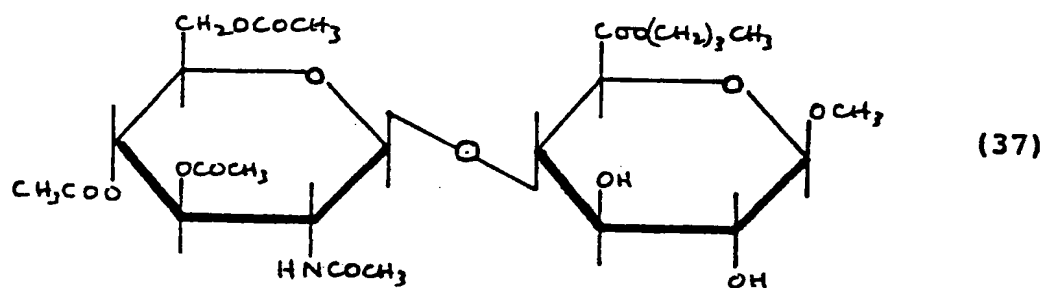
(36)

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( ii) hydrogenating the benzylated  $\beta$  -1,4 disaccharide to remove the benzyl groups to yield a  $\beta$  -1,4 partially acetylated disaccharide having the structure:

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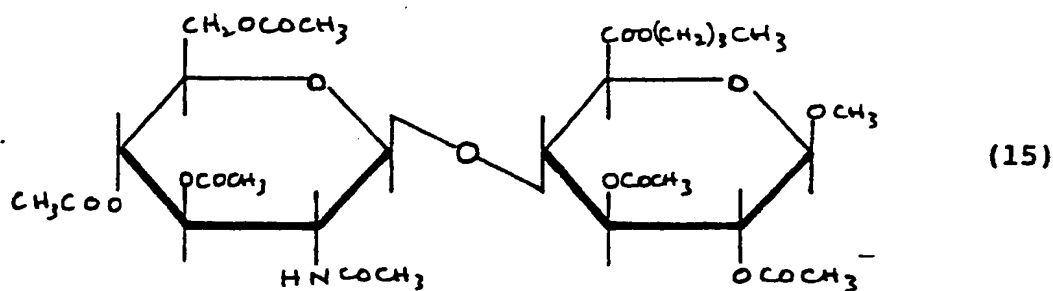


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and, (iii) acetylating the partially acetylated  $\beta$  -1,4 disaccharide to yield the fully acetylated  $\beta$  -1,4 disaccharide having the structure:

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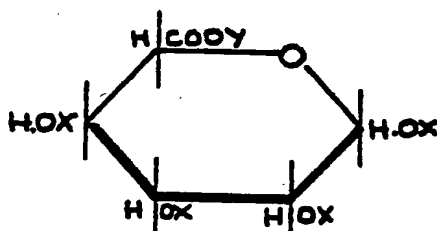


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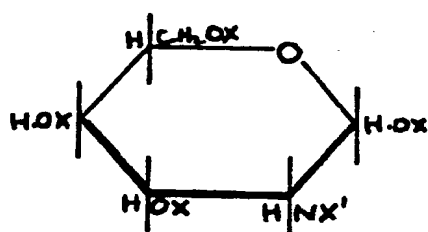
32. A composition suitable for topical application to mammalian skin comprising an effective amount of an esterified oligosaccharide as defined in any of claims 1 to 23, together with a cosmetically or physiologically acceptable carrier.

33. A composition suitable for topical application to mammalian skin comprising an effective amount of an esterified disaccharide including a uronic acid residue having the structure:



(50)

and a hexosamine residue having the structure:



(51)

where X is - H, C<sub>1</sub> to C<sub>4</sub> alkyl, -CO(CH<sub>2</sub>)<sub>m</sub>CH<sub>3</sub>, -SO<sub>3</sub>M,  
or an aryl radical,

X is - H, -CO(CH<sub>2</sub>)<sub>m</sub>CH<sub>3</sub>, or - SO<sub>3</sub>M,

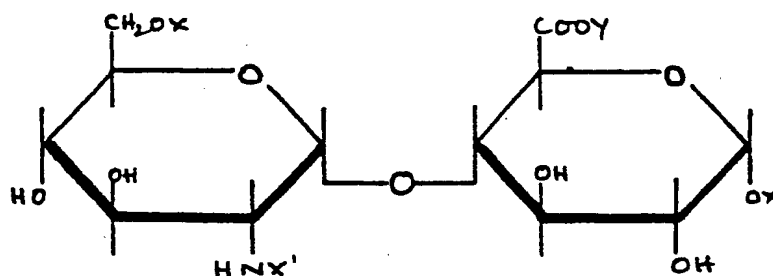
Y is - H, C<sub>1</sub> to C<sub>4</sub> alkyl or M, and

M is - H or a metallic or organic  
cation, and

m is 0 or the integer 1 or 2;

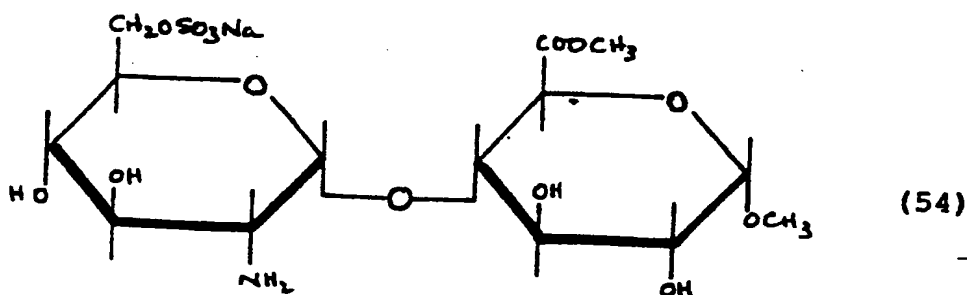
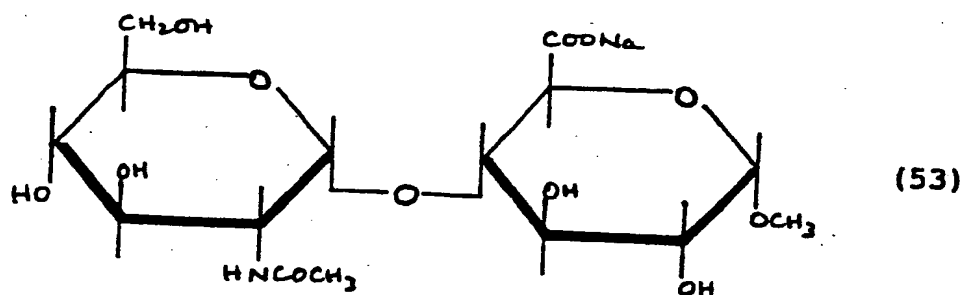
the groups designated X being the same or  
different, one -OH group from each pyranose ring  
structure being linked by a glycosidic linkage  
having the configuration  $\alpha$ -1,3,  $\beta$ -1,3,  
 $\alpha$ -1,4 or  $\beta$ -1,4; and the -COOY, -CH<sub>2</sub>OX and  
-OX groups being of either configuration with  
respect to the pyranose ring.

34. A composition according to claim 33 in which the  
esterified disaccharide has the structure:



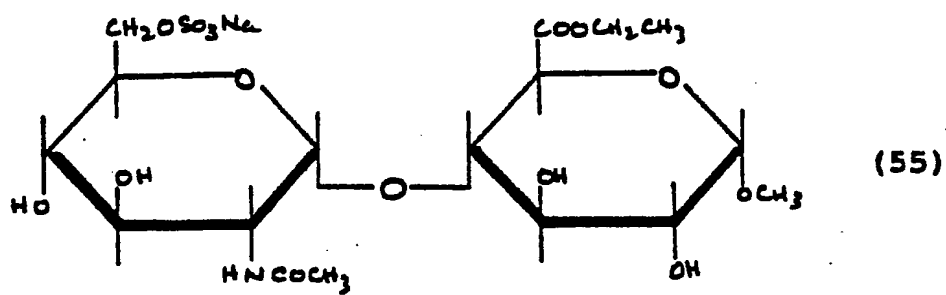
(52)

35. A composition according to claim 34, in which the esterified disaccharide is chosen from those having the following structures:



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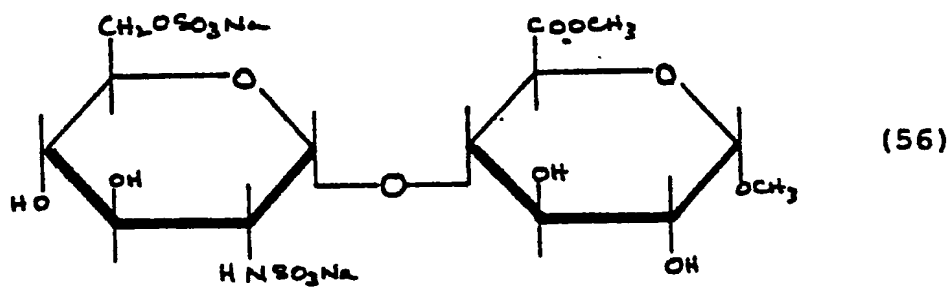
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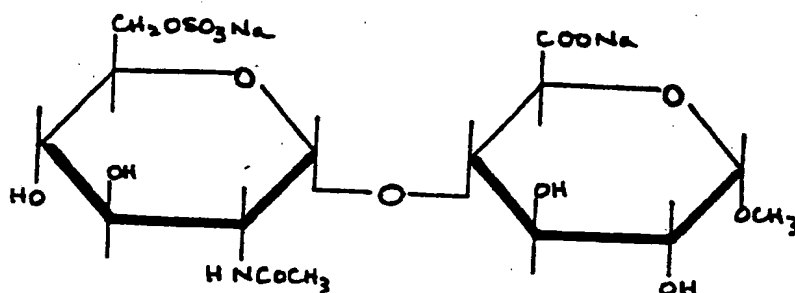
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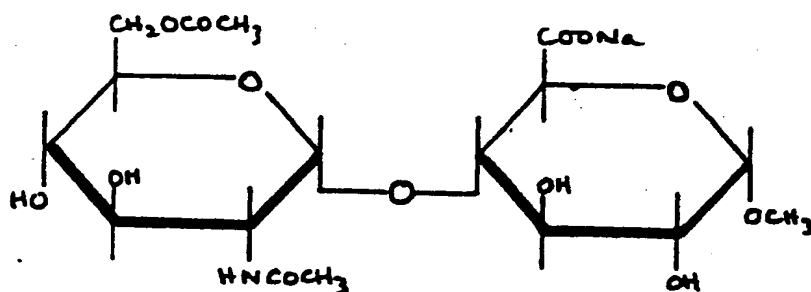
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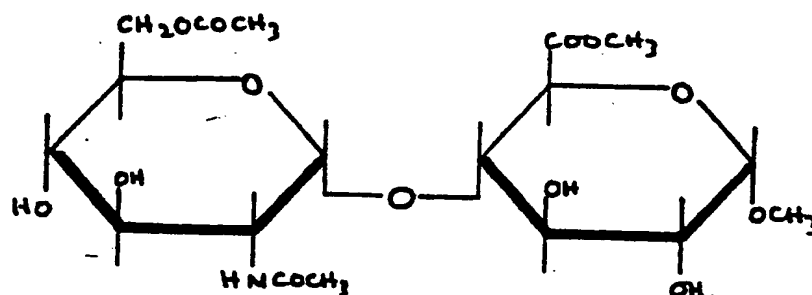
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(58)

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(59)

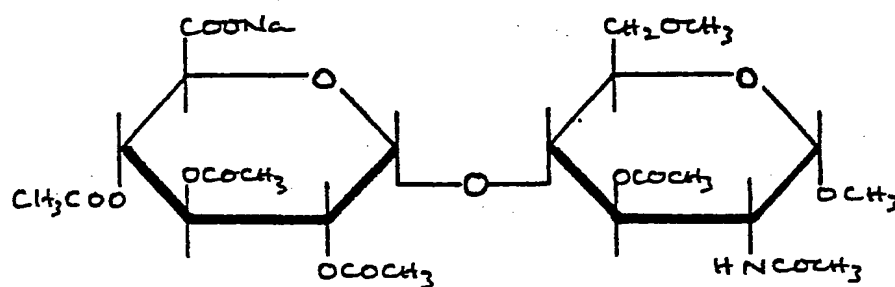
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36. A composition according to claim 33, in which the esterified disaccharide is chosen from those having the following structures:

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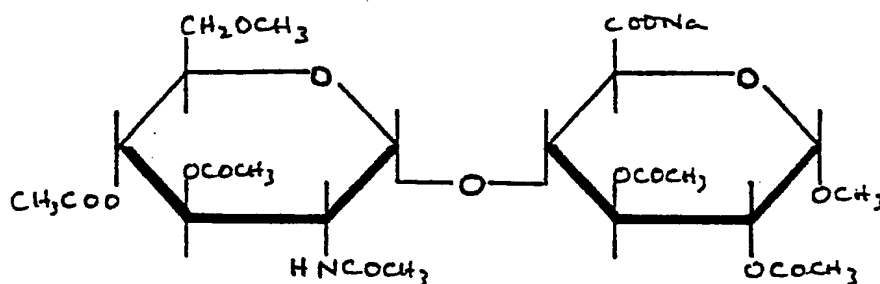
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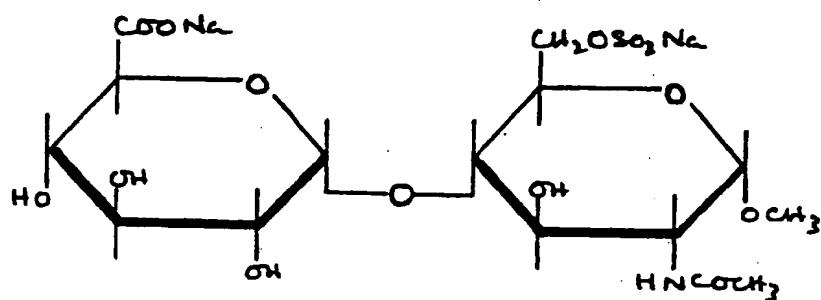
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(61)

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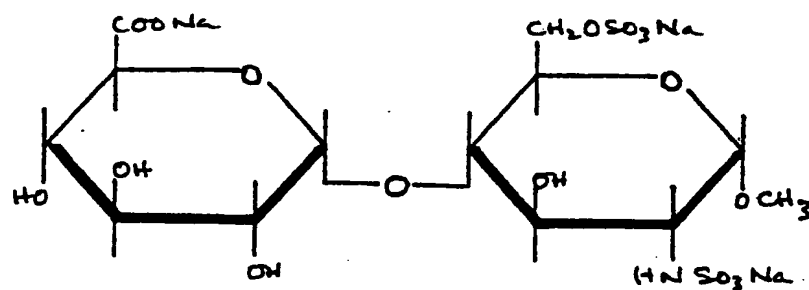
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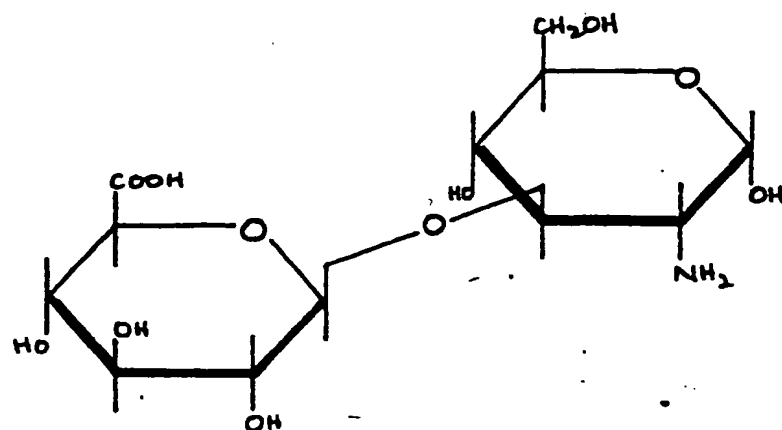
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(63)

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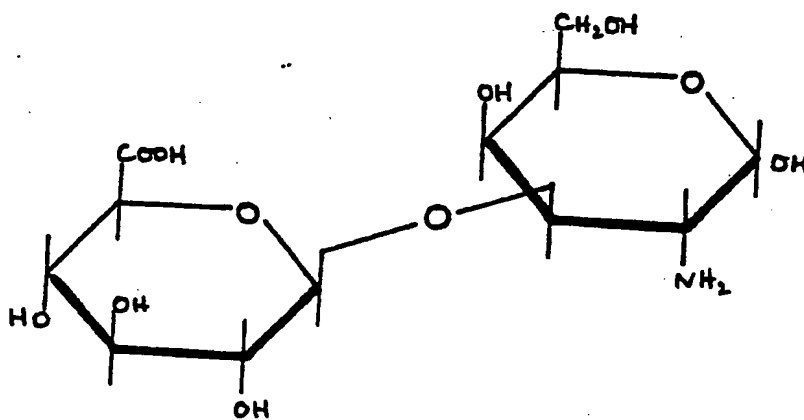
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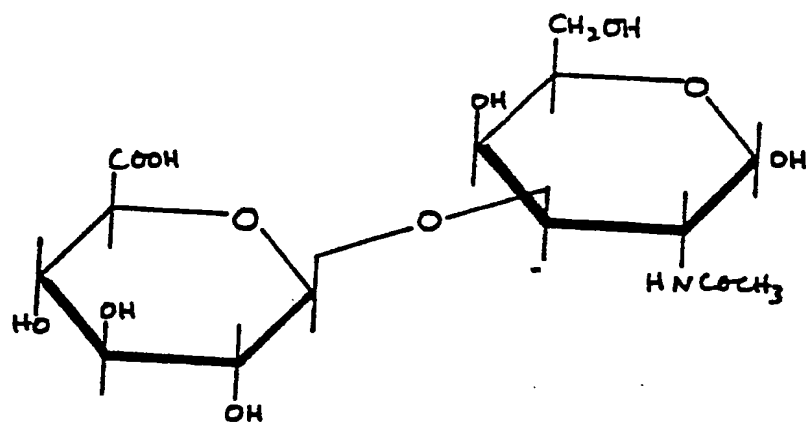
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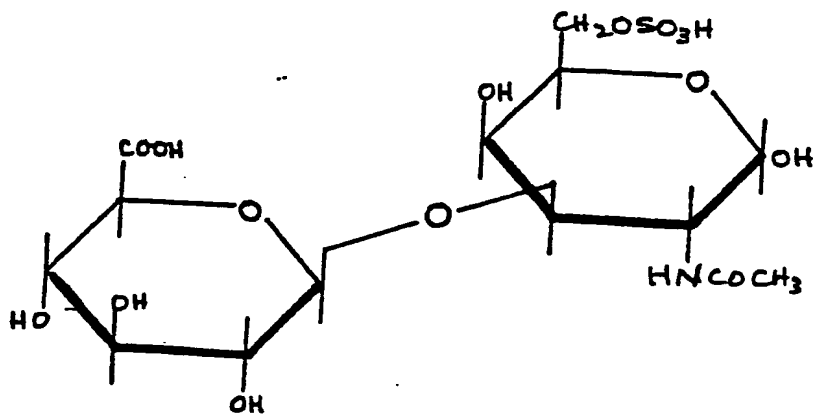




(30)



(31)



(32)

37. A composition suitable for topical application to mammalian skin comprising an effective amount of a low molecular weight limit enzymic digest comprising an esterified oligosaccharide derived from a glycosaminoglycan chosen from heparan sulphate, hyaluronic acid and chondroitin sulphate.
38. A composition according to any of claims 32 to 37, in which the effective amount of the esterified oligosaccharide present in the composition is from 0.01 to 10% by weight.
39. A composition according to any of claims 32 to 38, in which the effective amount of the esterified oligosaccharide present in the composition is from 0.01 to 5% by weight.
40. A composition according to any of claims 32 to 39, in which the carrier forms from 10 to 99.99% by weight.
41. A composition according to any of claims 32 to 40, in which the carrier forms from 50 to 99.5% by weight.
42. A composition according to any of claims 32 to 41, which additionally comprises a perfume.
43. A composition according to any of claims 32 to 42, which additionally comprises an activity enhancer.
44. A composition according to any of claims 32 to 43, which is in the form of a cream, a gel, a lotion or a stick.

45. A method for the conversion of vellus hair to growth as terminal hair, which comprises applying to mammalian skin in the region of vellus hair, an effective amount of a composition according to any of claims 32 to 44.

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46. A method for increasing the rate of terminal hair growth in mammalian species, which comprises applying to mammalian skin in the region of terminal hair, an effective amount of a composition according to any of claims 32 to 44.

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47. The use of an esterified oligosaccharide as defined in any of claims 1 to 31 in the promotion of mammalian hair growth or regrowth.

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48. The use of an esterified oligosaccharide as defined in any of claims 1 to 31 in the preparation of a composition for the treatment of baldness.

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